

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
7 February 2002 (07.02.2002)

PCT

(10) International Publication Number  
**WO 02/09768 A2**

(51) International Patent Classification?: **A61K 47/48**

(21) International Application Number: **PCT/US01/23747**

(22) International Filing Date: 27 July 2001 (27.07.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:  
60/220,707 27 July 2000 (27.07.2000) US  
60/261,337 12 January 2001 (12.01.2001) US

(71) Applicant (for all designated States except US): **RUTGERS, THE STATE UNIVERSITY [US/US]**; Old Queens Building, Somerset and George Streets, New Brunswick, NJ 08903 (US).

(71) Applicant and  
(72) Inventor: **UHRICH, Kathryn, E. [US/US]**; 920 Bloomfield Street, #1, Hoboken, NJ 07030 (US).

(74) Agent: **VIKSNINS, Ann, S.; Schwegman, Lundberg, Woessner & Kluth, P.O. Box 2938, Minneapolis, MN 55402 (US)**.

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SI, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.



**WO 02/09768 A2**

(54) Title: **THERAPEUTIC POLYESTERS AND POLYAMIDES**

(57) Abstract: Polymers (i.e. polyesters, polyamides, and polythioesters or a mixture thereof) which degrade hydrolytically into biologically active compounds are provided. Methods of producing these polymers, intermediates useful for preparing these polymers, and methods of using these polymers to deliver biologically active compounds to a host are also provided.

## THERAPEUTIC POLYESTERS AND POLYAMIDES

5

### Priority of Invention

This application claims priority from United States Provisional Application Number 60/220,707, filed 27 July 2000 and United States Provisional Application Number 60/261,337, filed 1 December 2001, which are incorporated herein by reference..

10

### Background of the Invention

Polyesters are used routinely by those skilled in the art in various drug delivery systems.

For example, U.S. Patent 5,942,252 describes a microcapsule comprising 15 as its biocompatible excipient a poly(lactide-co-glycolide), poly(lactide), poly(glycolide), copolyoxalate, polycaprolactone, poly(lactide-co-caprolactone), poly(esteramide), polyorthoester, poly(p-hydroxybutyric) acid and/or polyanhydride for use in delivering antigens or vaccines into and through mucosally-associated lymphoid tissue.

20 WO 99/29885 describes a process for degrading poly(ester-amides) and poly(ester-urethanes) encapsulating chemicals, drugs, enzymes, microorganisms and seeds by introducing the polymer to an aqueous nutrient solution and inoculating the solution with a culture containing a selected bacteria.

25 WO 98/36013 describes aliphatic-aromatic dihydroxy compounds for use as controlled drug delivery systems.

WO 97/39738 describes preparation of microparticles of a sustained release ionic conjugate comprising a free carboxyl group containing biodegradable polymers and a free amino group-containing drug.

30

### Summary of the Invention

Polyesters, polythioesters, and polyamides which degrade into useful biologically active compounds have now been developed. Accordingly, the invention provides a polymer of the invention which is polymer comprising a

backbone, wherein the backbone comprises ester, thioester, or amide linkages, and wherein the backbone comprises one or more groups that will yield a biologically active compound upon hydrolysis of the polymer.

The invention also provides a pharmaceutical composition comprising a 5 polymer of the invention and a pharmaceutically acceptable carrier.

The invention also provides a therapeutic method for treating a disease in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of the invention.

The invention also provides a method of delivering a biologically active 10 compound to a host comprising administering to the host a biocompatible and biodegradable polymer of the invention, which degrades into the biologically active compound.

The invention provides a polymer of the invention for use in medical 15 therapy, as well as the use of a polymer of the invention for the manufacture of a medicament useful for the treatment of a disease in a mammal, such as a human.

The invention also provides processes and intermediates disclosed herein that are useful for preparing a polymer of the invention.

#### Detailed Description of the Invention

##### 20 Definitions

The following definitions are used, unless otherwise described: halo is fluoro, chloro, bromo, or iodo. Alkyl, alkoxy, etc. denote both straight and branched groups; but reference to an individual radical such as "propyl" embraces only the straight chain radical, a branched chain isomer such as 25 "isopropyl" being specifically referred to. Aryl denotes a phenyl radical or an ortho-fused bicyclic carbocyclic radical having about nine to ten ring atoms in which at least one ring is aromatic. Heteroaryl encompasses a radical attached via a ring carbon of a monocyclic aromatic ring containing five or six ring atoms consisting of carbon and one to four heteroatoms each selected from the group 30 consisting of non-peroxide oxygen, sulfur, and N(X) wherein X is absent or is H, O, (C<sub>1</sub>-C<sub>6</sub>)alkyl, phenyl or benzyl, as well as a radical of an ortho-fused bicyclic heterocycle of about eight to ten ring atoms derived therefrom, particularly a

benz-derivative or one derived by fusing a propylene, trimethylene, or tetramethylene diradical thereto.

The term ester linkage means  $-OC(=O)-$  or  $-C(=O)O-$ ; the term thioester linkage means  $-SC(=O)-$  or  $-C(=O)S-$ ; and the term amide linkage means  $-$

5  $N(R)C(=O)-$  or  $-C(=O)N(R)-$ , wherein each R is a suitable organic radical, such as, for example, hydrogen,  $(C_1-C_6)alkyl$ ,  $(C_3-C_6)cycloalkyl$ ,  $(C_3-C_6)cycloalkyl(C_1-C_6)alkyl$ , aryl, heteroaryl, aryl $(C_1-C_6)alkyl$ , or heteroaryl $(C_1-C_6)alkyl$ .

The term "amino acid," comprises the residues of the natural amino acids

10 (e.g. Ala, Arg, Asn, Asp, Cys, Glu, Gln, Gly, His, Ile, Leu, Lys, Met, Phe, Pro, Ser, Thr, Trp, Tyr, and Val) in D or L form, as well as unnatural amino acids (e.g. phosphoserine, phosphothreonine, phosphotyrosine, hydroxyproline, gamma-carboxyglutamate; hippuric acid, octahydroindole-2-carboxylic acid, statine, 1,2,3,4,-tetrahydroisoquinoline-3-carboxylic acid, penicillamine, 15 ornithine, citruline,  $\alpha$ -methyl-alanine, para-benzoylphenylalanine, phenylglycine, propargylglycine, sarcosine, and tert-butylglycine). The term also comprises natural and unnatural amino acids bearing a conventional amino protecting group (e.g. acetyl or benzyloxycarbonyl), as well as natural and unnatural amino acids protected at the carboxy terminus (e.g. as a  $(C_1-C_6)alkyl$ , 20 phenyl or benzyl ester or amide; or as an  $\alpha$ -methylbenzyl amide). Other suitable amino and carboxy protecting groups are known to those skilled in the art (See for example, Greene, T.W.; Wutz, P.G.M. "Protecting Groups In Organic Synthesis" second edition, 1991, New York, John Wiley & sons, Inc., and references cited therein).

25 The term "host" includes animals and plants.

The term "peptide" describes a sequence of 2 to 35 amino acids (e.g. as defined hereinabove) or peptidyl residues. The sequence may be linear or cyclic.

For example, a cyclic peptide can be prepared or may result from the formation of disulfide bridges between two cysteine residues in a sequence. Preferably a

30 peptide comprises 3 to 20, or 5 to 15 amino acids. Peptide derivatives can be prepared as disclosed in U.S. Patent Numbers 4,612,302; 4,853,371; and 4,684,620, or as described in the Examples hereinbelow. Peptide sequences

specifically recited herein are written with the amino terminus on the left and the carboxy terminus on the right.

Polymers of the Invention

5        The biocompatible, biodegradable polyesters, polythioesters, and polyamides of the invention are useful in a variety of applications where delivery of a biologically active compound is desired. Examples of such applications include, but are not limited to, medical, dental and cosmetic uses.

10      The polymers of the invention may be prepared in accordance with methods commonly employed in the field of synthetic polymers to produce a variety of useful products with valuable physical and chemical properties. The polymers can be readily processed into pastes or solvent cast to yield films, coatings, microspheres and fibers with different geometric shapes for design of various medical implants, and may also be processed by compression molding 15 and extrusion.

Polyesters and polyamides prepared in accordance with the present invention have average molecular weights of about 1500 Daltons up to about 100,000 Daltons, calculated by Gel Permeation Chromatography (GPC) relative to narrow molecular weight polystyrene standards. Preferred polyesters and 20 polyamides have average molecular weights of about 1500 Daltons, up to about 50,000 Daltons calculated by Gel Permeation Chromatography (GPC) relative to narrow molecular weight polystyrene standards. Preferred polyesters and polyamides have average molecular weights of about 1500 Daltons, up to about 35,000 Daltons.

25      Medical implant applications include the use of polyesters, polythioesters, or polyamides to form shaped articles such as vascular grafts and stents, bone plates, sutures, implantable sensors, implantable drug delivery devices, stents for tissue regeneration, and other articles that decompose into non-toxic components within a known time period.

30      Polymers of the present invention can also be incorporated into oral formulations and into products such as skin moisturizers, cleansers, pads, plasters, lotions, creams, gels, ointments, solutions, shampoos, tanning products and lipsticks for topical application.

Although the invention provides homopolymers that are prepared from suitably functionalized biologically active compounds, Applicant has discovered that the mechanical and hydrolytic properties of polymers comprising one or more biologically active compounds can be controlled by incorporating a linking 5 group (L) into the polymer backbone.

Preferably, the polymers of the invention comprise backbones wherein biologically active compounds and linker groups are bonded together through ester linkages, thioester linkages, amide linkages, or a mixture thereof. Due to the presence of the ester, thioester, and/or amide linkages, the polymers can be 10 hydrolyzed under physiological conditions to provide the biologically active compounds. Thus, the polymers of the invention can be particularly useful as a controlled release source for a biologically active compound, or as a medium for the localized delivery of a biologically active compound to a selected site. For example, the polymers of the invention can be used for the localized delivery of 15 a therapeutic agent to a selected site within the body of a human patient (i.e. within or near a tumor), where the degradation of the polymer provides localized, controlled, release of the therapeutic agent.

#### Biologically Active Compounds

20 The term "biologically active compound" includes therapeutic agents that provide a therapeutically desirable effect when administered to an animal (e.g. a mammal, such as a human). Biologically active compounds that can be incorporated into the polymers of the invention possess at least two functional groups that can each be incorporated into an ester, thioester, or amide linkage of 25 a polymer (as discussed in detail below), such that, upon hydrolysis of the polymer, the therapeutic agent is obtained. These groups can independently be a hydroxy group (-OH), a mercapto group (-SH), an amine group (-NHR), or a carboxylic acid (-COOH).

The biologically active compounds can also comprise other functional 30 groups (including hydroxy groups, mercapto groups, amine groups, and carboxylic acids, as well as others) that can be used to modify the properties of the polymer (e.g. for branching, for cross linking, for appending other molecules (e.g. another biologically active compound) to the polymer, for changing the

solubility of the polymer, or for effecting the biodistribution of the polymer).

Lists of therapeutic agents can be found, for example, in: Physicians' Desk Reference, 55 ed., 2001, Medical Economics Company, Inc., Montvale, New Jersey; USPN Dictionary of USAN and International Drug Names, 2000, The

5 United States Pharmacopeial Convention, Inc., Rockville, Maryland; and The Merck Index, 12 ed., 1996, Merck & Co., Inc., Whitehouse Station, New Jersey. One skilled in the art can readily select therapeutic agents that possess the necessary functional groups for incorporation into the polymers of the invention from these lists.

10 Therapeutic agents that can be incorporated into the polymers of the invention include suitably functionalized analgesics or general or local anesthetics, anti-convulsants, anti-diabetic agents, anti-fibrotic agents, anti-infectives, anti-bacterials, anti-fungals, anti-neoplastics, cardioprotective agents, cardiovascular agents, anti-thrombotics, central nervous system stimulants, 15 cholinesterase inhibitors, contraceptives, deodorants, dopamine receptor agonists, erectile dysfunction agents, fertility agents, gastrointestinal agents, gout agents, hormones, immunomodulators, immunosuppressives, migraine agents, non-steroidal anti-inflammatory drugs (NSAIDs), motion sickness agents, muscle relaxants, nucleoside analogs, neurodegenerative agents (e.g., Parkinson's 20 disease), obesity agents, ophthalmic agents, osteoporosis agents, parasympatholytics, parasympathometrics, anti-anesthetics, prostaglandins, psychotherapeutic agents, respiratory agents, sedatives, hypnotics, skin and mucous membrane agents, smoking cessation agents, sympatholytics, urinary tract agents, vaginal agents, and vasodilators (see Physicians' Desk Reference, 25 55 ed., 2001, Medical Economics Company, Inc., Montvale, New Jersey, pages 201-202).

#### Linking Group "L"

The nature of the linking group "L" in a polymer of the invention is not 30 critical provided the polymer of the invention possesses acceptable mechanical properties and release kinetics for the selected therapeutic application. The linking group L is typically a divalent organic radical having a molecular weight

of from about 25 daltons to about 400 daltons. More preferably, L has a molecular weight of from about 40 daltons to about 200 daltons.

The linking group L typically has a length of from about 5 angstroms to about 100 angstroms using standard bond lengths and angles. More preferably,

- 5 the linking group L has a length of from about 10 angstroms to about 50 angstroms.

The linking group may be biologically inactive, or may itself possess biological activity. The linking group can also comprise other functional groups (including hydroxy groups, mercapto groups, amine groups, carboxylic acids, as 10 well as others) that can be used to modify the properties of the polymer (e.g. for branching, for cross linking, for appending other molecules (e.g. another biologically active compound) to the polymer, for changing the solubility of the polymer, or for effecting the biodistribution of the polymer).

15 **Specific And Preferred Values**

Specific and preferred values listed herein for radicals, substituents, groups, and ranges, are for illustration only; they do not exclude other defined values or other values within defined ranges for the radicals and substituents.

Specifically, (C<sub>1</sub>-C<sub>6</sub>)alkyl can be methyl, ethyl, propyl, isopropyl, butyl, 20 iso-butyl, sec-butyl, pentyl, 3-pentyl, or hexyl; (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl can be cyclopropyl, cyclobutyl, cyclopentyl, or cyclohexyl; (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl(C<sub>1</sub>-C<sub>6</sub>)alkyl can be cyclopropylmethyl, cyclobutylmethyl, cyclopentylmethyl, cyclohexylmethyl, 2-cyclopropylethyl, 2-cyclobutylethyl, 2-cyclopentylethyl, or 2-cyclohexylethyl; (C<sub>1</sub>-C<sub>6</sub>)alkoxy can be methoxy, ethoxy, propoxy, isopropoxy, 25 butoxy, iso-butoxy, sec-butoxy, pentoxy, 3-pentoxy, or hexyloxy; (C<sub>1</sub>-C<sub>6</sub>)alkanoyl can be acetyl, propanoyl or butanoyl; (C<sub>1</sub>-C<sub>6</sub>)alkoxycarbonyl can be methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, butoxycarbonyl, pentoxy carbonyl, or hexyloxycarbonyl; (C<sub>1</sub>-C<sub>6</sub>)alkylthio can be methylthio, ethylthio, propylthio, isopropylthio, butylthio, isobutylthio, 30 pentylthio, or hexylthio; (C<sub>2</sub>-C<sub>6</sub>)alkanoyloxy can be acetoxy, propanoyloxy, butanoyloxy, isobutanoyloxy, pentanoyloxy, or hexanoyloxy; aryl can be phenyl, indenyl, or naphthyl; and heteroaryl can be furyl, imidazolyl, triazolyl, triazinyl, oxazoyl, isoxazoyl, thiazoyl, isothiazoyl, pyrazoyl, pyrrolyl, pyrazinyl,

tetrazolyl, pyridyl, (or its N-oxide), thienyl, pyrimidinyl (or its N-oxide), indolyl, isoquinolyl (or its N-oxide) or quinolyl (or its N-oxide).

A specific biologically active compound that can be incorporated into the polymers of the invention is atorvastatin; enalapril; ranitidine; ciprofloxacin; 5 pravastatin; clarithromycin; cyclosporin; famotidine; leuprolide; acyclovir; paclitaxel; azithromycin; lamivudine; budesonide; albuterol; indinavir; metformin; alendronate; nizatidine; zidovudine; carboplatin; metoprolol; amoxicillin; diclofenac; lisinopril; ceftriaxone; captopril; salmeterol; xinafoate; imipenem; cilastatin; benazepril; cefaclor; ceftazidime; morphine; dopamine; 10 bialamicol; fluvastatin; phenamidine; podophyllinic acid 2-ethylhydrazine; acriflavine; chloroazodin; arsphenamine; amicarbilide; aminoquinuride; quinapril; oxymorphone; buprenorphine; butorphanol; nalbuphine. streptozocin; doxorubicin; daunorubicin; plicamycin; idarubicin; mitomycin C; pentostatin; mitoxantrone; cytarabine; fludarabine phosphate; floxuridine; cladribine; 6- 15 mercaptopurine; thioguanine; capecitabine; docetaxel; etoposide; gemcitabine; topotecan; vinorelbine; vincristine; vinblastine; teniposide; melphalan; methotrexate; 2-p-sulfanilylanilinoethanol; 4,4'-sulfinylidianiline; 4-sulfanilamidosalicylic acid; acediasulfone; acetosulfone; amikacin; amphotericin B; ampicillin; apalcillin; apiclycline; apramycin; arbekacin; aspoxicillin; 20 azidamfenicol; aztreonam; bacitracin; bambermycin(s); biapenem; brodimoprim; butirosin; capreomycin; carbenicillin; carbomycin; carumonam; cefadroxil; cefamandole; cefatrizine; cefbuperazone; cefclidin; cefdinir; cefditoren; cefepime; cefetamet; cefixime; cefmenoxime; cefminox; cefodizime; cefonicid; cefoperazone; ceforanide; cefotaxime; cefotetan; cefotiam; cefozopran; 25 cefpimizole; cefpiramide; cefpirome; cefprozil; cefroxadine; ceftetan; ceftibuten; cefuzonam; cephalexin; cephaloglycin; cephalosporin C; cephadrine; chloramphenicol; chlortetracycline; clinafloxacin; clindamycin; clomocycline; colistin; cyclacillin; dapsone; demeclocycline; diathymosulfone; dibekacin; dihydrostreptomycin; dirithromycin; doxycycline; enoxacin; eniromycin; 30 epicillin; erythromycin; flomoxef; fortimicin(s); gentamicin(s); glucosulfone solasulfone; gramicidin S; gramicidin(s); grepafloxacin; guamecycline; hetacillin; isepamicin; josamycin; kanamycin(s); leucomycin(s); lincomycin; lomefloxacin; lucensomycin; lymecycline; mecloxycline; meropenem;

methacycline; micronomicin; midecamycin(s); minocycline; moxalactam;  
mupirocin; nadifloxacin; natamycin; neomycin; netilmicin; norfloxacin;  
oleandomycin; oxytetracycline; p-sulfanilylbenzylamine; panipenem;  
paromomycin; pazufloxacin; penicillin N; pipacycline; pipemidic acid;  
5 polymyxin; primycin; quinacillin; ribostamycin; rifamide; rifampin; rifamycin  
SV; rifapentine; rifaximin; ristocetin; ritipenem; rokitamycin; rolitetracycline;  
rosaramycin; roxithromycin; salazosulfadimidine; sencycline; sisomicin;  
sparfloxacin; spectinomycin; spiramycin; streptomycin; succisulfone;  
sulfachrysoidine; sulfaloxic acid; sulfamidochrysoidine; sulfanilic acid;  
10 sulfoxone; teicoplanin; temafloxacin; temocillin; tetroxoprim; thiamphenicol;  
thiazolsulfone; thiostrepton; ticarcillin; tigemonam; tobramycin; tosufloxacin;  
trimethoprim; trospectomycin; trovafloxacin; tuberactinomycin; vancomycin;  
azaserine; candicidin(s); chlorphenesin; dermostatin(s); filipin; fungichromin;  
meparticin; nystatin; oligomycin(s); perimycin A; tubercidin; 6-azauridine; 6-  
15 diazo-5-oxo-L-norleucine; aclacinomycin(s); ancitabine; anthramycin;  
azacitadine; azaserine; bleomycin(s); carubicin; carzinophillin A; chlorozotocin;  
chromomycin(s); denopterin; doxifluridine; edatrexate; eflornithine; elliptinium;  
enocitabine; epirubicin; mannomustine; menogaril; mitobronitol; mitolactol;  
mopidamol; mycophenolic acid; nogalamycin; olivomycin(s); peplomycin;  
20 pirarubicin; piritrexim; prednimustine; procarbazine; pteropterin; puromycin;  
ranimustine; streptonigrin; thiamiprime; Tomudex® (N-[[5-[[1,4-Dihydro-2-  
methyl-4-oxo-6-quinazolinyl)methyl]methylamino]-2-thienyl]carbonyl]-L-  
glutamic acid), trimetrexate, tubercidin, ubenimex, vindesine, zorubicin;  
argatroban; coumetarol; dicoumarol; ethyl biscoumacetate; ethylidene  
25 dicoumarol; iloprost; lamifiban; taprostene; tioclomarol; tirofiban; amiprilose;  
bucillamine; gusperimus; mycophenolic acid; procodazole; romurtide; sirolimus  
(rapamycin); tacrolimus; butethamine; fenalcomine; hydroxytetracaine;  
naepaine; orthocaine; piridocaine; salicyl alcohol; 3-amino-4-hydroxybutyric  
acid; aceclofenac; alminoprofen; amfenac; bromfenac; bromosaligenin;  
30 bumadizon; carprofen; diclofenac; diflunisal; ditazol; enfenamic acid; etodolac;  
etofenamate; fendosal; fepradinol; flufenamic acid; gentisic acid; glucamethacin;  
glycol salicylate; meclofenamic acid; mefenamic acid; mesalamine; niflumic

acid; olsalazine; oxaceprol; S-adenosylmethionine; salicylic acid; salsalate; sulfasalazine; or tolfenamic acid.

A preferred biologically active compound suitable for incorporation into polyesters of the invention is morphine, dopamine, bialamicol, or tetracycline.

5 A preferred biologically active compound suitable for incorporation into polyamides of the present invention is phenamidine, acriflavine, chloroazodin, arsphenamine, amicarbilide or aminoquinuride.

Another preferred biologically active compound that can be incorporated into a polymer of the invention is oxymorphone, buprenorphine, butorphanol, or 10 nalbuphine.

Another preferred biologically active compound that can be incorporated into a polymer of the invention is methotrexate, doxorubicin, or daunorubicin.

Another preferred biologically active compound that can be incorporated into a polymer of the invention is atorvastatin, enalapril, ranitidine, pravastatin,

15 cyclosporin, famotidine, leuprolide, acyclovir, lamivudine, budesonide, albuterol, indinavir, metformin, alendronate, nizatidine, zidovudine, carboplatin, metoprolol, lisinopril, captopril, salmeterol, cilastatin, benazepril, cefaclor, fluvastatin, quinapril, gemcitabine or vincristine.

Another preferred biologically active compound that can be incorporated 20 into a polymer of the invention is a nonsteroidal anti-inflammatory drug, for example, a nonsteroidal anti-inflammatory drug as described in U.S. Patent Application (Serial Number 09/732,516, filed 07 December 2000), 3-amino-4-hydroxybutyric acid, aceclofenac, alminoprofen, amfenac, bromfenac, bromosaligenin, bumadizon, carprofen, diclofenac, diflunisal, ditazol, enfenamic 25 acid, etodolac, etofenamate, fendosal, fepradinol, flufenamic acid, gentisic acid, glucamethacin, glycol salicylate, meclofenamic acid, mefenamic acid, mesalamine, niflumic acid, olsalazine, oxaceprol, S-adenosylmethionine, salicylic acid, salsalate, sulfasalazine, tolfenamic acid and the like.

Another preferred biologically active compound that can be incorporated 30 into a polymer of the invention is an anti-bacterial, for example, 2-p-sulfanilylanilinoethanol, 4,4'-sulfinyldianiline, 4-sulfanilamidosalicylic acid, acediasulfone, acetosulfone, amikacin, amoxicillin, amphotericin B, ampicillin, apalcillin, apicycline, apramycin, arbekacin, aspoxicillin, azidamfenicol,

azithromycin, aztreonam, bacitracin, bambemycin(s), biapenem, brodimoprim, butirosin, capreomycin, carbenicillin, carbomycin, carumonam, cefadroxil, cefamandole, cefatrizine, cefbuperazone, cefclidin, cefdinir, cefditoren, cefepime, cefetamet, cefixime, cefmenoxime, cefminox, cefodizime, cefonicid, 5 cefoperazone, ceforanide, cefotaxime, cefotetan, cefotiam, cefozopran, cefpimizole, cefpiramide, cefpirome, cefprozil, cefroxadine, ceftazidime, cefteram, cefributen, ceftriaxone, cefuzonam, cephalexin, cephaloglycin, cephalosporin C, cephradine, chloramphenicol, chlortetracycline, ciprofloxacin, clarithromycin, clinafloxacin, clindamycin, clomocycline, colistin, cyclacillin, 10 dapsone, demeclocycline, diathymosulfone, dibekacin, dihydrostreptomycin, dirithromycin, doxycycline, enoxacin, eniromycin, epicillin, erythromycin, flomoxef, fortimicin(s), gentamicin(s), glucosulfone solasulfone, gramicidin S, gramicidin(s), grepafloxacin, guamecycline, hetacillin, imipenem, isepamicin, josamycin, kanamycin(s), leucomycin(s), lincomycin, lomefloxacin, 15 lucensomycin, lymecycline, meclocycline, meropenem, methacycline, micromycin, midecamycin(s), minocycline, moxalactam, mupirocin, nadifloxacin, natamycin, neomycin, netilmicin, norfloxacin, oleandomycin, oxytetracycline, p-sulfanilylbenzylamine, panipenem, paromomycin, pazufloxacin, penicillin N, pipacycline, pipemidic acid, polymyxin, primycin, 20 quinacillin, ribostamycin, rifamide, rifampin, rifamycin SV, rifapentine, rifaximin, ristocetin, ritipenem, rokitamycin, rolitetracycline, rosaramycin, roxithromycin, salazosulfadimidine, sancycline, sisomicin, sparfloxacin, spectinomycin, spiramycin, streptomycin, succisulfone, sulfachrysoidine, sulfaloxic acid, sulfamidochrysoidine, sulfanilic acid, sulfoxone, teicoplanin, 25 temafloxacin, temocillin, tetracycline, tetroxoprim, thiamphenicol, thiazolsulfone, thiostrepton, ticarcillin, tigemonam, tobramycin, tosufloxacin, trimethoprim, trospectomycin, trovafloxacin, tuberactinomycin, vancomycin and the like.

Another preferred biologically active compound that can be incorporated

30 into a polymer of the invention is an anti-fungal, for example, azaserine, candidin(s), chlorphenesin, dermostatin(s), filipin, fungichromin, meparticin, nystatin, oligomycin(s), perimycin A, tubercidin and the like.

Another preferred biologically active compound that can be incorporated into a polymer of the invention is an anti-cancer (e.g., carcinomas, sarcomas, leukemias and cancers derived from cells of the nervous system), including anti-neoplastic, for example, 6-azauridine, 6-diazo-5-oxo-L-norleucine, 6-

5 mercaptoperine, aclacinomycin(s), ancitabine, anthramycin, azacitidine, azaserine, bleomycin(s), capecitabine, carubicin, carzinophillin A, chlorozotocin, chromomycin(s), cladribine, cytarabine, daunorubicin, denopterin, docetaxel, doxifluridine, doxorubicin, edatrexate, eflornithine, elliptinium, enocitabine, epirubicin, etoposide, floxuridine, fludarabine, gemcitabine, idarubicin,

10 mannomustine, melphalan, menogaril, methotrexate, mitobronitol, mitolactol, mitomycin C, mitoxantrone, mopidamol, mycophenolic acid, nogalamycin, olivomycin(s), paclitaxel, pentostatin, peplomycin, pirarubicin, piritrexim, plicamycin, podophyllinic acid 2-ethylhydrazine, prednimustine, procarbazine, pteropterin, puromycin, ranimustine, streptonigrin, streptozocin, teniposide,

15 thiamiprime, thioguanine, Tomudex® (N-[[5-[[1,4-Dihydro-2-methyl-4-oxo-6-quinazolinyl)methyl]methylamino]-2-thienyl]carbonyl]-L-glutamic acid), toptecan, trimetrexate, tubercidin, ubenimex, vinblastine, vindesine, vinorelbine, zorubicin and the like.

Another preferred biologically active compound that can be incorporated into a polymer of the invention is an anti-thrombotic, for example, argatroban, coumetarol, dicoumarol, ethyl biscoumacetate, ethylidene dicoumarol, iloprost, lamifiban, taprostene, tioclomarol, tirofiban and the like.

Another preferred biologically active compound that can be incorporated into a polymer of the invention is an immunosuppressive, for example, 6-

25 mercaptoperine, amiprilose, bucillamine, gusperimus, mycophenolic acid, procodazole, romurtide, sirolimus (rapamycin), tacrolimus, ubenimex and the like.

Another preferred biologically active compound that can be incorporated into a polymer of the invention is a general or local anesthetic, for example,

30 butethamine, fenalcomine, hydroxytetracaine, naepaine, orthocaine, piridocaine, salicyl alcohol and the like.

A specific value for L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 1 to 25 carbon atoms, wherein one

or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-), and wherein the chain is optionally substituted on carbon with one or more (e.g. 1, 2, 3, or 4) substituents selected from the group consisting of (C<sub>1</sub>-C<sub>6</sub>)alkoxy, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyloxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxycarbonyl, (C<sub>1</sub>-C<sub>6</sub>)alkylthio, azido, cyano, nitro, halo, hydroxy, oxo (=O), carboxy, aryl, aryloxy, heteroaryl, and heteroaryloxy.

Another specific value for L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 1 to 25 carbon atoms, wherein the chain is optionally substituted on carbon with one or more (e.g. 1, 2, 3, or 4) substituents selected from the group consisting of (C<sub>1</sub>-C<sub>6</sub>)alkoxy, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyloxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxycarbonyl, (C<sub>1</sub>-C<sub>6</sub>)alkylthio, azido, cyano, nitro, halo, hydroxy, oxo, carboxy, aryl, aryloxy, heteroaryl, and heteroaryloxy.

Another specific value for L is an amino acid.

Another specific value for L is a peptide

Another specific value for L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 1 to 25 carbon atoms, wherein one or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-).

A more specific value for L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 3 to 15 carbon atoms, wherein one or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-), and wherein the chain is optionally substituted on carbon with one or more (e.g. 1, 2, 3, or 4) substituents selected from the group consisting of (C<sub>1</sub>-C<sub>6</sub>)alkoxy, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyloxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxycarbonyl, (C<sub>1</sub>-C<sub>6</sub>)alkylthio, azido, cyano, nitro, halo, hydroxy, oxo, carboxy, aryl, aryloxy, heteroaryl, and heteroaryloxy.

Another more specific value for L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 3 to 15 carbon atoms, wherein one or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-).

Another more specific value for L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 3 to 15 carbon atoms.

Another more specific value for L is a divalent, branched or unbranched, hydrocarbon chain, having from 3 to 15 carbon atoms.

A preferred value for L is a divalent, branched or unbranched, hydrocarbon chain, having from 6 to 10 carbon atoms.

5 A more preferred value for L is a divalent hydrocarbon chain having 7, 8, or 9 carbon atoms.

A most preferred value for L is a divalent hydrocarbon chain having 8 carbon atoms.

10 A specific polymer of the invention comprises one or more units of formula (I):



wherein  $R_1$  is group that will provide a biologically active compound upon hydrolysis of the polymer; each A is independently an amide linkage, a thioester linkage, or an ester linkage; and L is a linking group.

15 Another specific polymer of the invention is a polymer which comprises one or more units of formula (II) in the backbone:



20 wherein:  $R_2$  and  $R_3$  are each independently a group that will yield a biologically active compound upon hydrolysis of the polymer; each A is independently an amide, thioester, or ester linkage; and each L is independently a linking group. Such a polymer, wherein  $R_2$  and  $R_3$  are groups that will yield differing biologically active compounds upon hydrolysis of the polymer, are particularly 25 useful for the administration of a combination of two therapeutic agents to an animal.

A preferred group of polyesters and polyamides includes polymers that are comprised of compounds containing at least two free alcohol or phenol groups or two at least two free amine groups available for reactions which 30 co-polymerize with carboxylic acid groups or bis(acyl) chlorides.

### Formulations

The polymers of the invention can be formulated as pharmaceutical compositions and administered to a mammalian host, such as a human patient in a variety of forms adapted to the chosen route of administration, i.e., orally, rectally, or parenterally, by intravenous, intramuscular, intraperitoneal, 5 intraspinal, intracranial, topical or subcutaneous routes. For some routes of administration, the polymer can conveniently be formulated as micronized particles.

Thus, the present compounds may be systemically administered, e.g., orally, in combination with a pharmaceutically acceptable vehicle such as an 10 inert diluent or an assimilable edible carrier. They may be enclosed in hard or soft shell gelatin capsules, may be compressed into tablets, or may be incorporated directly with the food of the patient's diet. For oral therapeutic administration, the active compound may be combined with one or more excipients and used in the form of ingestible tablets, buccal tablets, troches, 15 capsules, elixirs, suspensions, syrups, wafers, and the like. Such compositions and preparations preferably contain at least 0.1% of polymer by weight. The percentage of the compositions and preparations may, of course, be varied and may conveniently be between about 2 to about 80% of the weight and preferably 2 to about 60 % of a given unit dosage form. The amount of polymer in such 20 therapeutically useful compositions is such that an effective dosage level will be obtained.

The tablets, troches, pills, capsules, and the like may also contain the following: binders such as gum tragacanth, acacia, corn starch or gelatin; excipients such as dicalcium phosphate; a disintegrating agent such as corn 25 starch, potato starch, alginic acid and the like; a lubricant such as magnesium stearate; and a sweetening agent such as sucrose, fructose, lactose or aspartame or a flavoring agent such as peppermint, oil of wintergreen, or cherry flavoring may be added. When the unit dosage form is a capsule, it may contain, in addition to materials of the above type, a liquid carrier, such as a vegetable oil or 30 a polyethylene glycol. Various other materials may be present as coatings or to otherwise modify the physical form of the solid unit dosage form. For instance, tablets, pills, or capsules may be coated with gelatin, wax, shellac or sugar and the like. A syrup or elixir may contain the active compound, sucrose or fructose

as a sweetening agent, methyl and propylparabens as preservatives, a dye and flavoring such as cherry or orange flavor. Of course, any material used in preparing any unit dosage form should be pharmaceutically acceptable and substantially non-toxic in the amounts employed. In addition, the active 5 compound may be incorporated into sustained-release preparations and devices.

The polymer may also be administered intravenously, intraspinally, intracranially, or intraperitoneally by infusion or injection. Solutions of the polymer can be prepared a suitable solvent such as an alcohol, optionally mixed with a nontoxic surfactant. Dispersions can also be prepared in glycerol, liquid 10 polyethylene glycols, triacetin, and mixtures thereof and in oils. Under ordinary conditions of storage and use, these preparations contain a preservative to prevent the growth of microorganisms.

The pharmaceutical dosage forms suitable for injection or infusion can include sterile solutions or dispersions or sterile powders comprising the 15 polymer containing the active ingredient which are adapted for the extemporaneous preparation of sterile injectable or infusible solutions or dispersions, optionally encapsulated in liposomes. In all cases, the ultimate dosage form should be sterile, fluid and stable under the conditions of manufacture and storage. The liquid carrier or vehicle can be a solvent or liquid 20 dispersion medium comprising, for example, ethanol, a polyol (for example, glycerol, propylene glycol, liquid polyethylene glycols, and the like), vegetable oils, nontoxic glyceryl esters, and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the formation of liposomes, by the maintenance of the required particle size in the case of dispersions or by the use 25 of surfactants. The prevention of the action of microorganisms can be brought about by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars, buffers or sodium chloride. Prolonged absorption of the injectable compositions can be 30 brought about by the use in the compositions of agents delaying absorption, for example, aluminum monostearate and gelatin.

Sterile injectable solutions are prepared by incorporating the polymer in the required amount in the appropriate solvent with various of the other

5 ingredients enumerated above, as required, followed by filter sterilization. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and freeze drying techniques, which yield a powder of the active ingredient plus any additional desired ingredient present in the previously sterile-filtered solutions.

For topical administration, the present polymers can be applied in pure form. However, it will generally be desirable to administer them as compositions or formulations, in combination with a dermatologically acceptable carrier, which may be a solid or a liquid.

10 Useful solid carriers include finely divided solids such as talc, clay, microcrystalline cellulose, silica, alumina and the like. Useful liquid carriers include water, alcohols or glycols or water-alcohol/glycol blends, in which the present compounds can be dissolved or dispersed at effective levels, optionally with the aid of non-toxic surfactants. Adjuvants such as fragrances and  
15 additional antimicrobial agents can be added to optimize the properties for a given use. The resultant liquid compositions can be applied from absorbent pads, used to impregnate bandages and other dressings, or sprayed onto the affected area using pump-type or aerosol sprayers.

Thickeners such as synthetic polymers, fatty acids, fatty acid salts and  
20 esters, fatty alcohols, modified celluloses or modified mineral materials can also be employed with liquid carriers to form spreadable pastes, gels, ointments, soaps, and the like, for application directly to the skin of the user.

#### Dosages

25 Useful dosages of the polymers can be determined by comparing their *in vitro* activity, and *in vivo* activity of the therapeutic agent in animal models. Methods for the extrapolation of effective dosages in mice, and other animals, to humans are known to the art; for example, see U.S. Pat. No. 4,938,949. Additionally, useful dosages can be determined by measuring the rate of  
30 hydrolysis for a given polymer under various physiological conditions. The amount of a polymer required for use in treatment will vary not only with the particular polymer selected but also with the route of administration, the nature

of the condition being treated and the age and condition of the patient and will be ultimately at the discretion of the attendant physician or clinician.

The desired dose may conveniently be presented in a single dose or as divided doses administered at appropriate intervals, for example, as two, three,

5 four or more sub-doses per day. The sub-dose itself may be further divided, e.g., into a number of discrete loosely spaced administrations.

### Combination Therapies

The polymers of the invention are also useful for administering a

10 combination of therapeutic agents to an animal. Such a combination therapy can be carried out in the following ways: 1) a second therapeutic agent can be dispersed within the polymer matrix of a polymer of the invention, and can be released upon degradation of the polymer; 2) a second therapeutic agent can be appended to a polymer of the invention (i.e. not in the backbone of the polymer)

15 with bonds that hydrolyze to release the second therapeutic agent under physiological conditions; 3) the polymer of the invention can incorporate two therapeutic agents into the polymer backbone (e.g. a polymer comprising one or more units of formula (II)) or 4) two polymers of the invention, each with a different therapeutic agent can be administered together (or within a short period

20 of time).

Thus, the invention also provides a pharmaceutical composition comprising a polymer of the invention and a second therapeutic agent that is dispersed within the polymer matrix of a polymer of the invention. The invention also provides a pharmaceutical composition comprising a polymer of the invention having a second therapeutic agent appended to the polymer (e.g. with bonds that will hydrolyze to release the second therapeutic agent under physiological conditions).

The polymers of the invention can also be administered in combination with other therapeutic agents that are effective to treat a given condition to

30 provide a combination therapy. Thus, the invention also provides a method for treating a disease in a mammal comprising administering an effective amount of a combination of a polymer of the invention and another therapeutic agent. The invention also provides a pharmaceutical composition comprising a polymer of

the invention, another therapeutic agent, and a pharmaceutically acceptable carrier.

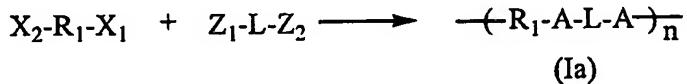
Preferred drug combinations for incorporation into the polymers or the compositions of the invention include the following: amoxicillin/clavulanic acid; and imipenem/cilastatin.

Preparation Of Polymers Of The Invention

Processes for preparing polymers of the invention are provided as further embodiments of the invention and are illustrated by the following procedures in which the meanings of the generic radicals are as given above unless otherwise qualified.

For example, a polymer of the invention can be prepared, as illustrated in Scheme I, from a biologically active compound of formula (X<sub>1</sub>-R<sub>1</sub>-X<sub>2</sub>) and a linker precursor of formula Z<sub>1</sub>-L-Z<sub>2</sub>, wherein X<sub>1</sub>, X<sub>2</sub>, Z<sub>1</sub>, and Z<sub>2</sub> are selected from the values in the table below.

Scheme I



The biologically active compound and the linker precursor can be polymerized using well known synthetic techniques (e.g. by condensation) to provide a

polymer of the invention (Ia) wherein each A is independently an ester linkage, a thioester linkage, or an amide linkage.

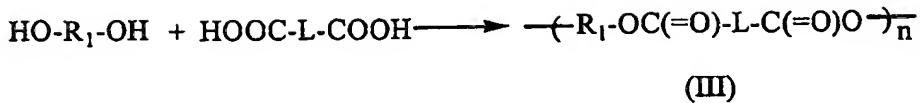
Depending on the reactive functional group (X<sub>1</sub> or X<sub>2</sub>) of the biologically active compound, a corresponding functional group (Z<sub>1</sub> or Z<sub>2</sub>) can be selected from the following table, to provide an ester linkage, thioester linkage, or amide linkage in the polymer backbone.

	Functional Group On Biologically active compound (X <sub>1</sub> or X <sub>2</sub> )	Functional Group On Linker Precursor (Z <sub>1</sub> or Z <sub>2</sub> )	Resulting Linkage In Polymer
5	-COOH	-OH	Ester
	-COOH	-NHR	Amide
	-COOH	-SH	Thioester
	-OH	-COOH	Ester
10	-SH	-COOH	Thioester
	-NHR	-COOH	Amide
	-SO <sub>3</sub> H	-OH	Sulfate Ester
	-OH	-SO <sub>3</sub> H	Sulfate Ester

As will be clear to one skilled in the art, suitable protecting groups can be used during the reaction illustrated in Scheme I (and in the reactions illustrated in Schemes II-XV below). For example, other functional groups present in the biologically active compound or the linker precursor can be protected during polymerization, and the protecting groups can subsequently be removed to provide the polymer of the invention. Suitable protecting groups and methods for their incorporation and removal are well known in the art (see for example Greene, T.W.; Wutz, P.G.M. "Protecting Groups In Organic Synthesis" second edition, 1991, New York, John Wiley & sons, Inc.).

Additionally, when a carboxylic acid is reacted with a hydroxy group, a mercapto group, or an amine group to provide an ester linkage, thioester linkage, or an amide linkage, the carboxylic acid can be activated prior to the reaction, for example, by formation of the corresponding acid chloride. Numerous methods for activating carboxylic acids, and for preparing ester linkages, thioester linkages, and amide linkages, are known in the art (see for example Advanced Organic Chemistry: Reaction Mechanisms and Structure, 4 ed., Jerry March, John Wiley & Sons, pages 419-437 and 1281).

A polyester of the invention can be formed from a biologically active compound of formula (HO-R<sub>1</sub>-OH) and from a linker precursor of formula HOOC-L-COOH as illustrated in Scheme II.

SCHEME II

Reaction of the hydroxy groups of the biologically active compound with the carboxylic acids of the linker precursor provides a polymer of formula (III), which is a polymer of the invention.

5        A preferred biologically active dihydroxy compound that can be used to prepare a polyester of the invention is: amikacin; amphotericin B; apicycline; apramycin; arbekacin; azidamfenicol; bambermycin(s); butirosin, carbomycin; cefpiramide; chloramphenicol; chlortetracycline; clindamycin; clomocycline; demeclocycline; diathymosulfone; dibekacin, dihydrostreptomycin;

10      dirithromycin; doxycycline; erythromycin; fortimicin(s); gentamicin(s); glucosulfone solasulfone; guamecycline; isepamicin; josamycin; kanamycin(s); leucomycin(s); lincomycin; lucensomycin; lymecycline; mecloxycline; methacycline; micromomicin; midecamycin(s); minocycline; mupirocin; natamycin; neomycin; netilmicin; oleandomycin; oxytetracycline; paromomycin;

15      pipacycline; podophyllinic acid 2-ethylhydrazine; primycin; ribostamycin; rifamide; rifampin; rifamycin SV; rifapentine; rifaximin; ristocetin; rokitamycin; rolitetracycline; rosaramycin; roxithromycin; sancycline; sisomicin; spectinomycin; spiramycin; streptomycin; teicoplanin; tetracycline; thiampenicol; thiostrepton; tobramycin; trospectomycin; tuberactinomycin;

20      vancomycin; candicidin(s); chlorphenesin; dermostatin(s); filipin; fungichromin; meparticin; nystatin; oligomycin(s); perimycin A; tubercidin; 6-azauridine; aclacinomycin(s); ancitabine; anthramycin; azacitidine; bleomycin(s); carubicin; carzinophillin A; chlorozotocin; chromomycin(s); doxifluridine; enocitabine; epirubicin; gemcitabine; mannustine; menogaril; atorvastatin; pravastatin;

25      clarithromycin; leuprolide; paclitaxel; mitobronitol; mitolactol; mopidamol; nogalamycin; olivomycin(s); peplomycin; pirarubicin; prednimustine; puromycin; ranimustine; tubercidin; vindesine; zorubicin; coumetarol; dicoumarol; ethyl biscoumacetate; ethylidene dicoumarol; iloprost; taprostene; tioclofarol; amiprilose; romurtide; sirolimus (rapamycin); tacrolimus; salicyl

30      alcohol; bromosaligenin; ditazol; fepradolin; gentisic acid; glucamethacin;

olsalazine; S-adenosylmethionine; azithromycin; salmeterol; budesonide; albuterol; indinavir; fluvastatin; streptozocin; doxorubicin; daunorubicin; plicamycin; idarubicin; pentostatin; mitoxantrone; cytarabine; fludarabine phosphate; floxuridine; cladribine; capecitabine; docetaxel; etoposide; topotecan; 5 vinblastine; or teniposide.

A polyamide of the invention can be prepared using a procedure similar to that illustrated in Scheme II by replacing the biologically active dihydroxy compound in Scheme II with a suitable biologically active diamino compound. A preferred biologically active diamino compound that can be used to prepare a 10 polymer of the invention is: 2-p-sulfanilylanilinoethanol; 4,4'-sulfinyldianiline; acediasulfone; acetosulfone; amikacin; apramycin; arbekacin; bacitracin; brodimorprim; butirosin; colistin; capreomycin; dapsone; dibekacin; eniromycin; gramicidin S; polymyxin; teicoplanin; fortimicin(s); gentamicin(s); glucosulfone solasulfone; grepafloxacin; imipenem; isepamicin; kanamycin(s); 15 lymecycline; micromycin; neomycin; netilmicin; p-sulfanilylbenzylamine; paromomycin; ribostamycin; ristocetin; sisomicin; sparfloxacin; spectinomycin; sulfachrysoidine; sulfamidochrysoidine; sulfoxone; tetroxoprim; thiazolsulfone; tobramycin; trimethoprim; edatrexate; eflornithine; mannomustine; mitoxantrone; peplomycin; piritrexim; procarbazine; pteropterin; trimetrexate; 20 gusperimus; butethamine; naepaine; piridocaine; trospectomycin; tuberactinomycin; vancomycin; candidin(s); meparticin; perimycin A; ranitidine; famotidine; metformin; nizatidine; carboplatin; lisinopril; methotrexate; mitomycin bleomycin(s) or thioguanine.

A polythioester of the invention can be prepared using a procedure 25 similar to that illustrated in Scheme II by replacing the biologically active dihydroxy compound in Scheme II with a suitable biologically active dimercapto compound.

A polysulfate ester of the invention can be formed by replacing the 30 dicarboxylic acid linker compound with a disulfo acid compound

A polyester/polyamide of the invention can be formed from a biologically active compound of formula (HRN-R<sub>1</sub>-OH) and from a linker precursor of formula HOOC-L-COOH as illustrated in Scheme III.

SCHEME III

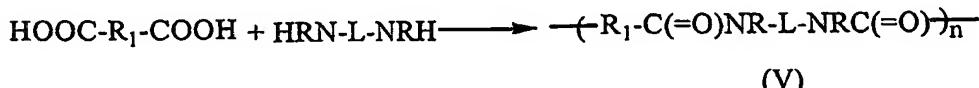
Reaction of the hydroxy group and the amino group of the biologically active compound with the carboxylic acids of the linker precursor provides a polymer of formula (IV), which is a polymer of the invention.

5        A preferred biologically active hydroxy/amino compound that can be used to prepare a polyester/polyamide of the invention is: 2-p-sulfanilylanilinoethanol; 4-sulfanilamidosalicylic acid; amikacin; amphotericin B; apramycin; arbekacin; aspoxicillin; butirosin; capreomycin; cefadroxil; cefatrizine; cefdinir; cefprozil; dibekacin; dihydrostreptomycin; dirithromycin; 10      enviomycin; gramicidin(s); teicoplanin; fortimicin(s); gentamicin(s); glucosulfone solasulfone; isepamicin; kanamycin(s); lucensomycin; lymecycline; meropenem; micromycin; natamycin; neomycin; netilmicin; paromomycin; ribostamycin; ristocetin; sisomicin; spectinomycin; streptomycin; thiostrepton; tobramycin; trospectomycin; tuberactinomycin; vancomycin; 15      candicidin(s); mepartycin; nystatin; perimycin A; tubercidin; anthramycin; azacitadine; bleomycin(s); carubicin; carcinophillin A; cytarabine; denopterin; elliptinium; epirubicin; gemcitabine; mannomustine; peplomycin; pirarubicin; pteropterin; puromycin; streptonigrin; tubercidin; ubenimex; vindesine; zorubicin; gusperimus; ubenimex; fenalcomine; hydroxytetracaine; orthocaine; 20      3-amino-4-hydroxybutyric acid; etofenamate; fepradolin; mesalamine; S-adenosylmethionine; leuprolide; acyclovir; paclitaxel; lamivudine; albuterol; indinavir; alendronate; zidovudine; metoprolol; amoxicillin; salmeterol; imipenem; doxorubicin; daunorubicin; idarubicin; pentostatin; mitoxantrone; fludarabine phosphate; floxuridine; cladribine; vinorelbine; vincristine; or 25      vinblastine.

A polythioester/polyamide of the invention can be prepared using a procedure similar to that illustrated in Scheme II by replacing the hydroxy/amino biologically active compound in Scheme II with a suitable mercapto/amino biologically active compound.

A polyamide of the invention can be formed from a biologically active compound of formula (HOOC-R<sub>1</sub>-COOH) and from a linker precursor of formula HRN-L-NRH as illustrated in Scheme IV.

SCHEME IV



5 Reaction of the carboxylic acid groups of the biologically active compound with the amino groups of the linker precursor provides a polymer of formula (V), which is a polymer of the invention.

A preferred biologically active dicarboxylic acid compound that can be used to prepare a polyamide of the invention is: bambermycin(s); carbenicillin;

10 carzinophillin A; cefixime; cefminox; cefpimizole; cefodizime; cefonicid; ceforanide; cefotetan; ceftibuten; cephalosporin C; denopterin; edatrexate; moxalactam; olsalazine; penicillin N; quinacillin; temocillin; ticarcillin; Tomudex® (N-[[5-[[1,4-Dihydro-2-methyl-4-oxo-6-quinazolinyl)methyl]methylamino]-2-thienyl]carbonyl]-L-glutamic acid);  
15 lisinopril; cilastatin; ceftazidime; or methotrexate.

A polyester of the invention can be prepared using a procedure similar to that illustrated in Scheme IV by replacing the diamino linker precursor with a dihydroxy linker precursor. Similarly, a polyester/polyamide of the invention can be prepared using a procedure similar to that illustrated in Scheme IV by

20 replacing the diamino linker precursor with an hydroxy/amino linker precursor; and a polythioester/polyamide of the invention can be prepared using a procedure similar to that illustrated in Scheme IV by replacing the diamino linker precursor with an mercapto/amino linker precursor.

A polyester of the invention can be formed from a biologically active compound of formula (HO-R<sub>1</sub>-COOH) and from a linker precursor of formula HO-L-COOH as illustrated in Scheme V.

SCHEME V

Reaction of the hydroxy group and the carboxylic acid of the biologically active



compound, with the carboxylic acid and the hydroxy group of the linker precursor provides a polymer of formula (VI), which is a polymer of the

5 invention.

A preferred biologically active hydroxy/carboxylic acid compound that can be used to prepare a polymer of the invention is: 4-sulfanilamidosalicylic acid; amphotericin B; apalcillin; apicycline; aspoxicillin; bambermycin(s); biapenem; cefadroxil; cefamandole; cefatrizine; cefbuperazone; cefdinir; 10 cefonicid; cefoperazone; cefpiramide; cefprozil; enviomycin; teicoplanin; flomoxef; glycol salicylate; lucensomycin; lymecycline; meropenem; moxalactam; mupirocin; nadifloxacin; natamycin; panipenem; podophyllinic acid 2-ethylhydrazine; ritipenem; salazosulfadimidine; sulfaloxic acid; vancomycin; 3-amino-4-hydroxybutyric acid; candididin(s); carzinophillin A; 15 denopterin; diflunisal; fendosal; gentisic acid; iloprost; lamifiban; mesalamine; nystatin; olsalazine; oxaceprol; pteropterin; romurtide; salicylic acid; salsalate; streptonigrin; sulfasalazine; taprostene; ubenimex; amoxicillin; pravastatin; imipenem; mycophenolic acid; or fluvastatin.

A polyester/polyamide of the invention can be prepared using a 20 procedure similar to that illustrated in Scheme V by replacing biologically active hydroxy/carboxylic compound with a biologically active amino/carboxylic acid compound. A preferred biologically active amino/carboxylic acid compound that can be used to prepare a polymer of the invention is: 3-amino-4-hydroxybutyric acid; 4-sulfanilamidosalicylic acid; 6-diazo-5-oxo-L-norleucine; 25 aceclofenac; acediasulfone; alminoprofen; amfenac; amphotericin B; ampicillin; argatroban; aspoxicillin; azaserine; aztreonam; bromfenac; bumadizon; candididin(s); carprofen; carumonam; carzinophillin A; cefadroxil; cefatrizine; cefclidin; cefdinir; cefditoren; cefepime; cefetamet; cefixime; cefmenoxime; cefminox; cefodizime; ceforanide; cefotaxime; cefotiam; cefozopran; cefpirome; 30 cefprozil; cefroxadine; ceftazidime; cefteram; cefributen; ceftriaxone;

cefuzonam; cephalexin; cephaloglycin; cephalosporin C; cephadrine; clinafloxacin; cyclacillin; denopterin; edatrexate; eflornithine; enfenamic acid; enoxacin; epicillin; etodolac; enviomycin; teicoplanin; flufenamic acid; grepafloxacin; hetacillin; imipenem; lomefloxacin; lucensomycin; lymecycline; 5 meclofenamic acid; mefenamic acid; meropenem; mesalamine; natamycin; niflumic acid; norfloxacin; nystatin; pazufloxacin; penicillin N; pipemidic acid; procodazole; pteropterin; S-adenosylmethionine; sparfloxacin; streptonigrin; succisulfone; sulfachrysoidine; temafloxacin; tigemonam; tirofiban; tolfenamic acid; tosufloxacin; trovafloxacin; ubenimex; vancomycin; enalapril; amoxicillin; 10 ciprofloxacin; diclofenac; lisinopril; ceftriaxone; cilastatin; benazepril; cefaclor; ceftazidime; quinapril; melphalan; or methotrexate.

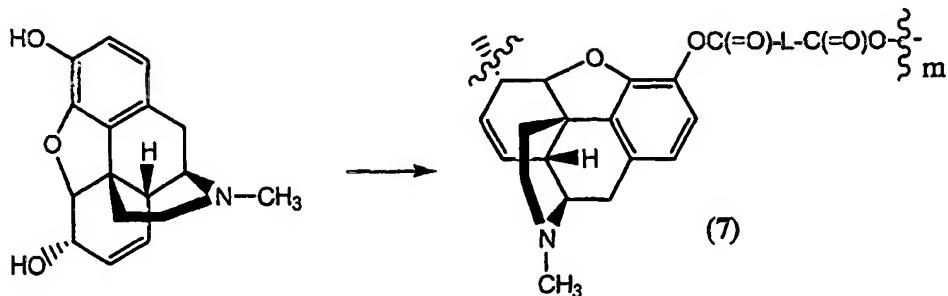
A polythioester/polyester of the invention can be prepared using a procedure similar to that illustrated in Scheme V by replacing biologically active hydroxy/carboxylic compound with a biologically active mercapto/carboxylic acid compound. A preferred biologically active mercapto/carboxylic acid compound that can be used to prepare a polymer of the invention is bucillamine or captopril.

A polysulfonamide of the invention can be prepared using a procedure similar that illustrated in Scheme V by replacing the biologically active 20 hydroxy/carboxylic compound with a biologically active amine/sulfo acid compound. A preferred biologically active amine/sulfo acid compound that can be used to prepare a polymer of the invention is: sulfanilic acid or sulfoxone.

In the polymers of formulae (I, and III-VI) illustrated in Schemes I-V above, R<sub>1</sub>, A, L, and R can have any of the values, specific values, or preferred 25 values described herein.

The co-polymerization of morphine with a diacid chloride to provide a polyester of the invention is depicted in Scheme VI.

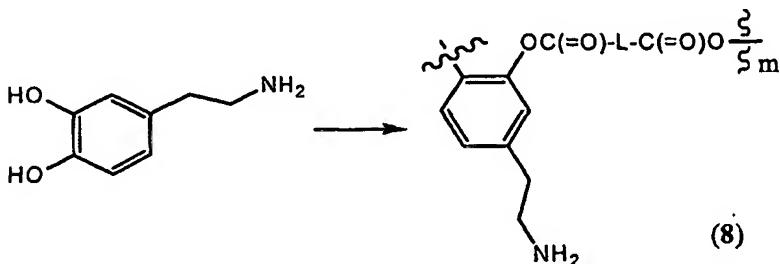
Scheme VI



In the reaction illustrated in Scheme VI, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula (7) wherein L has any of the values, specific values, or preferred values described herein is a 5 preferred polymer of the invention. For a polymer of formula (7), m is an integer that is greater than or equal to 2.

The co-polymerization of dopamine with bis(acyl) chloride to provide a polyester of the invention is depicted in Scheme VII.

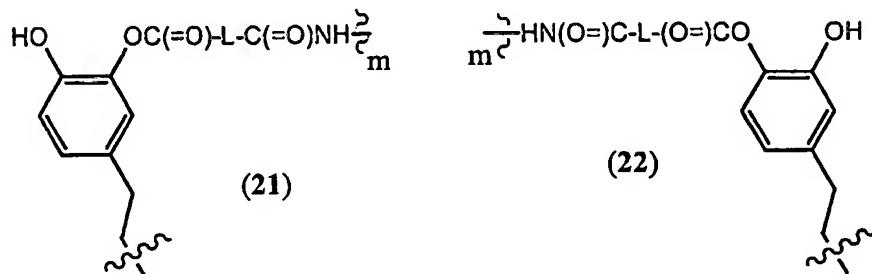
10

Scheme VII

In the reaction illustrated in Scheme VII, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula (8) wherein 15 L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (8), m is an integer that is greater than or equal to 2. Prior to the polymerization illustrated in Scheme VII, the amino group of dopamine can be protected with a suitable protecting group, which can subsequently be removed, to provide the polymer of 20 the invention.

It should be noted that dopamine can also be incorporated into a polyester/polyamide of the invention by reacting the amino group and either

hydroxy group of dopamine with a compound of formula HOOC-L-COOH, or an activated derivative thereof to provide a compound of formula (21) or (22):

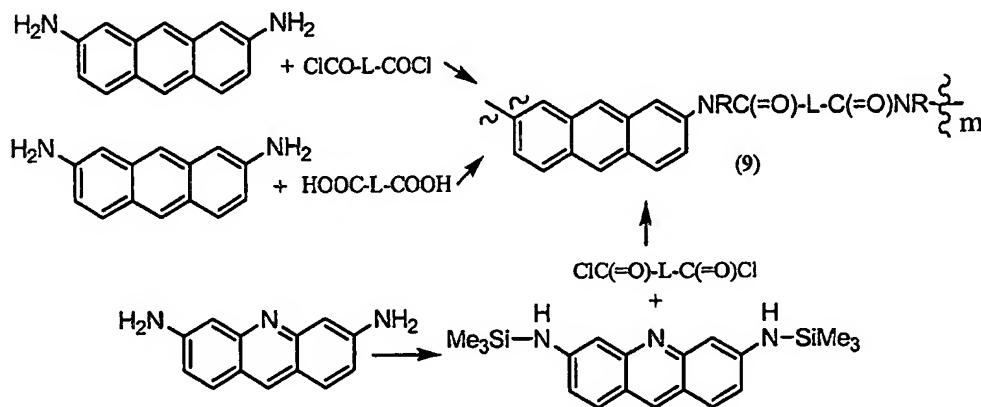


5 Prior to the polymerization, the hydroxy group which will not be polymerized can be protected with a suitable protecting group, which can subsequently be removed to provide the polymer of the invention.

The co-polymerization of acriflavine to provide a polyamide of the invention is depicted in Scheme VIII.

10

### Scheme VIII



The diamino groups of acriflavine are copolymerized in solution (preferably high-boiling point organic solvent such as dimethylformamide) with an activated dicarboxylic acid (e.g. sebacoyl chloride). The polyamide is isolated by methods

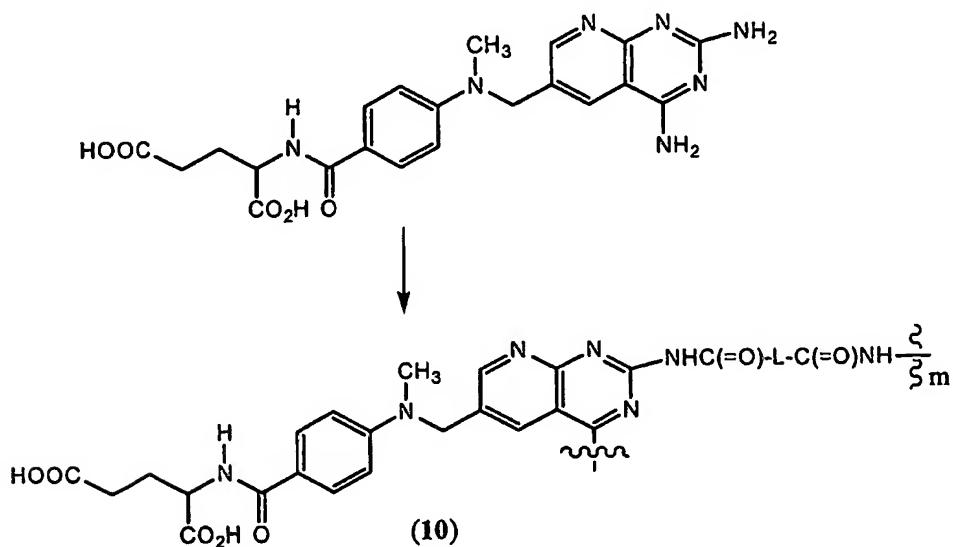
15 well known in the art. Alternatively, the amino groups can be reacted with a dicarboxylic acid by employing high temperatures (e.g. in the melting range), or a coupling agent. This process of making polyamides is also well known to those skilled in the art. In yet another embodiment, the diamino groups can be activated in the presence of hexamethylsilazine to form silylated amines. The

silylated amines can then be allowed to react with an activated dicarboxylic acid (e.g. sebacyl chloride) to provide a polymer of the invention.

In the reaction illustrated in Scheme VIII, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula 5 (9) wherein L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (9), m is an integer that is greater than or equal to 2.

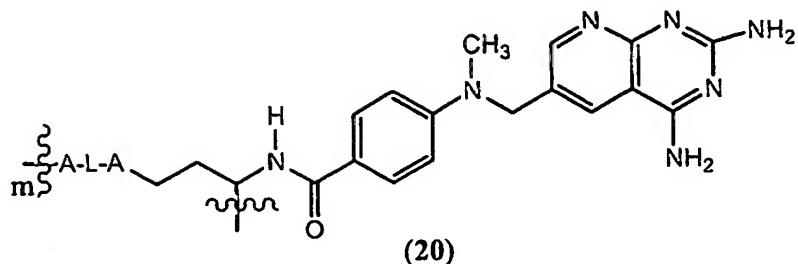
The preparation of a polymer of the invention comprising methotrexate is illustrated in Scheme IX.

10

Scheme IX

In the reaction illustrated in Scheme IX, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula (10) 15 wherein L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (10), m is an integer that is greater than or equal to 2. Prior to the polymerization illustrated in Scheme IX, the carboxylic acids of methotrexate can be protected with suitable protecting groups, which can subsequently be removed, to provide 20 the polymer of the invention.

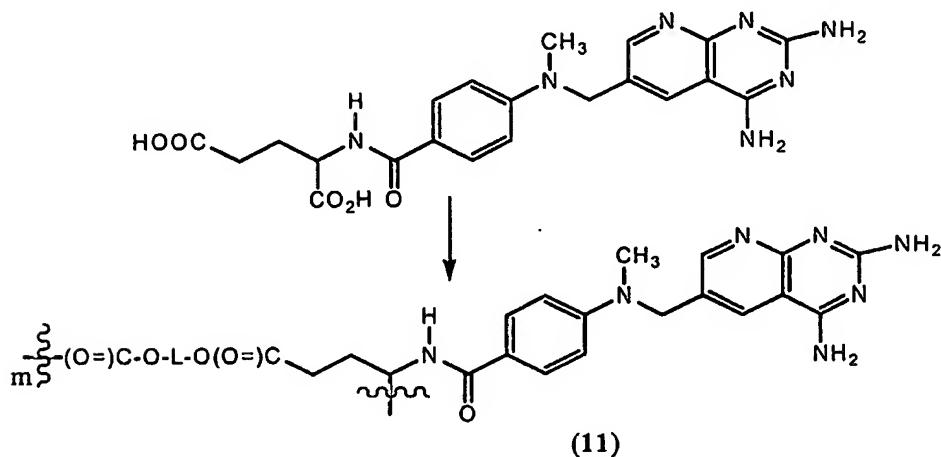
It will be appreciated by one skilled in the art that a polymer of the invention of the following formula (20):



wherein each A is independently an ester linkage, a thioester linkage, or an amide linkage can be prepared as illustrated in Scheme X by selecting a linker precursor with the appropriate functionality. For a compound of formula (20) the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A 5 polymer of formula (20) wherein L and each A has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (20), m is an integer that is greater than or equal to 2.

10 The preparation of another polymer of the invention comprising methotrexate is illustrated in Scheme X.

Scheme X

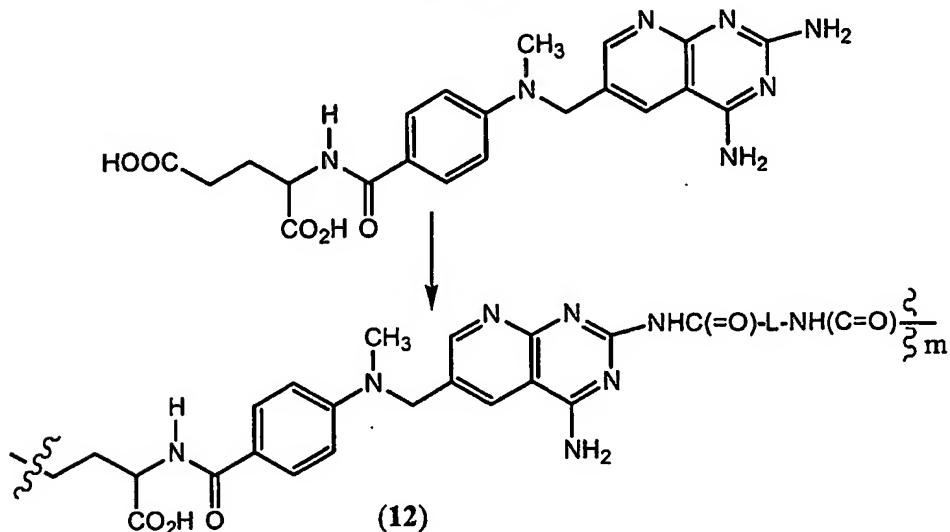


15 In the reaction illustrated in Scheme X, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula (11) wherein L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (11), m

is an integer that is greater than or equal to 2. Prior to the polymerization illustrated in Scheme X, the amino groups of methotrexate can be protected with suitable protecting groups, which can subsequently be removed, to provide the polymer of the invention.

5 The preparation of another polymer of the invention comprising methotrexate is illustrated in Scheme XI.

Scheme XI



10 In the reaction illustrated in Scheme XI, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_5-$ . A polymer of formula (12) wherein L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (12), m is an integer that is greater than or equal to 2. Prior to the polymerization illustrated in Scheme XI, the carboxylic acid and amino group of methotrexate that are not reacted to form the polymer can be protected with suitable protecting groups, which can subsequently be removed, to provide the polymer of the invention.

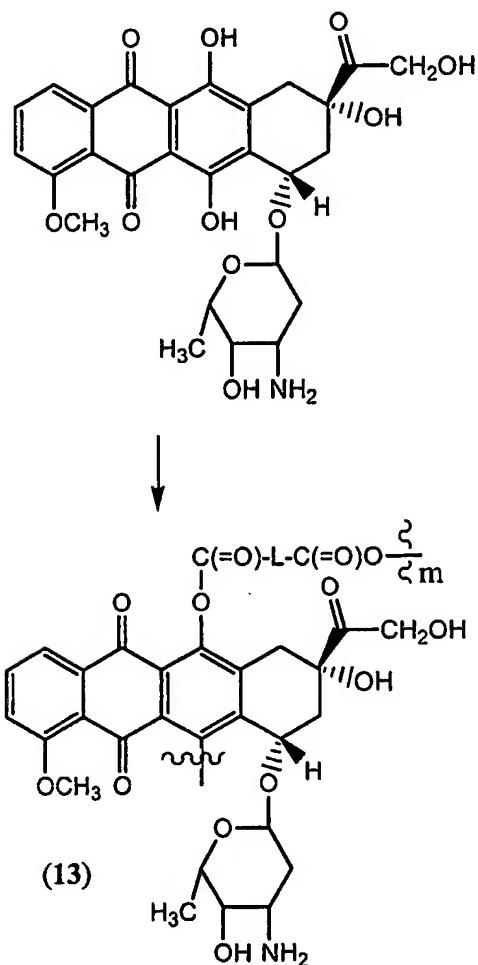
15

20 A polymer of the invention that comprises methotrexate is particularly useful for treating psoriasis, inflammatory bowel disease, skin cancer, or brain tumors. Such a polymer is also particularly useful as an anti-neoplastic agent

anti-infective agent, and for local administration of an anti-tumor agent following a lumpectomy or mastectomy.

The preparation of a polymer of the invention comprising doxorubicin is 5 illustrated in Scheme XII.

Scheme XII



10

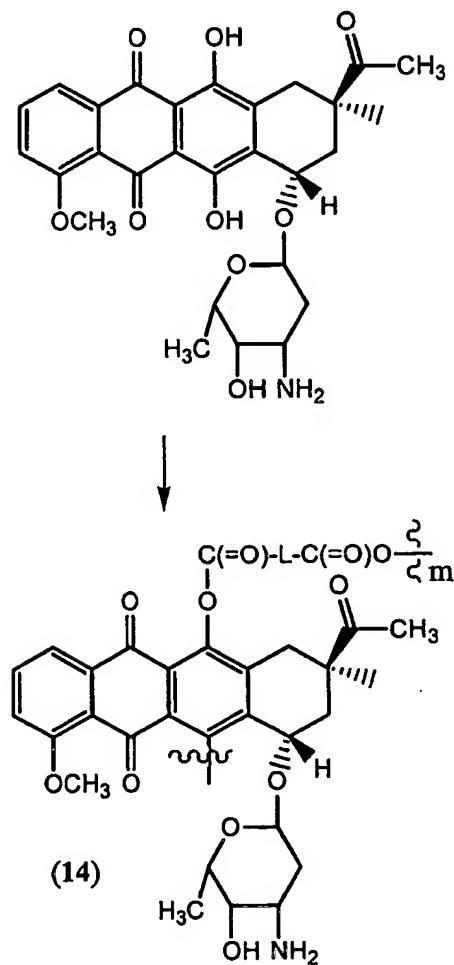
In the reaction illustrated in Scheme XII, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula (13) wherein L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of

formula (13), m is an integer that is greater than or equal to 2. Prior to the polymerization illustrated in Scheme XII, the functional groups of doxorubicin that are not reacted to form the polymer can be protected with suitable protecting groups, which can subsequently be removed, to provide the polymer of the 5 invention.

A polymer of the invention that comprises doxorubicin is particularly useful for local administration as an anti-tumor (e.g. brain tumor) agent or anti-neoplastic agent.

The preparation of a polymer of the invention comprising daunorubicin is 10 illustrated in Scheme XIII.

Scheme XIII



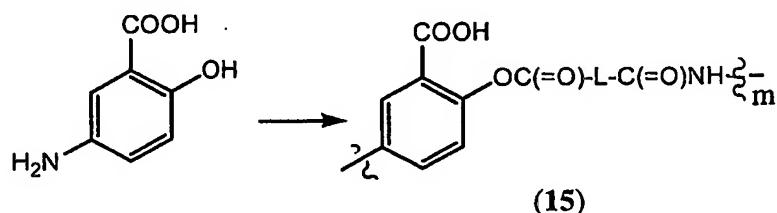
In the reaction illustrated in Scheme XIII, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula (14) wherein L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (14), m is an integer that is greater than or equal to 2. Prior to the polymerization illustrated in Scheme XIII, the functional groups of daunorubicin that are not reacted to form the polymer can be protected with suitable protecting groups, which can subsequently be removed, to provide the polymer of the invention.

10 A polymer of the invention that comprises daunorubicin is particularly useful for local administration as an anti-tumor (e.g. brain tumor) agent or an anti-neoplastic agent.

The preparation of a polymer of the invention comprising 5-aminosalicylic acid is illustrated in Scheme XIV.

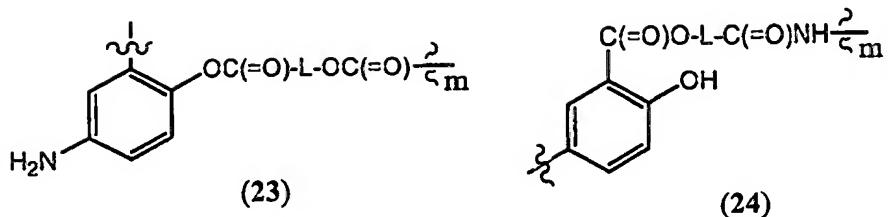
15

Scheme XIV



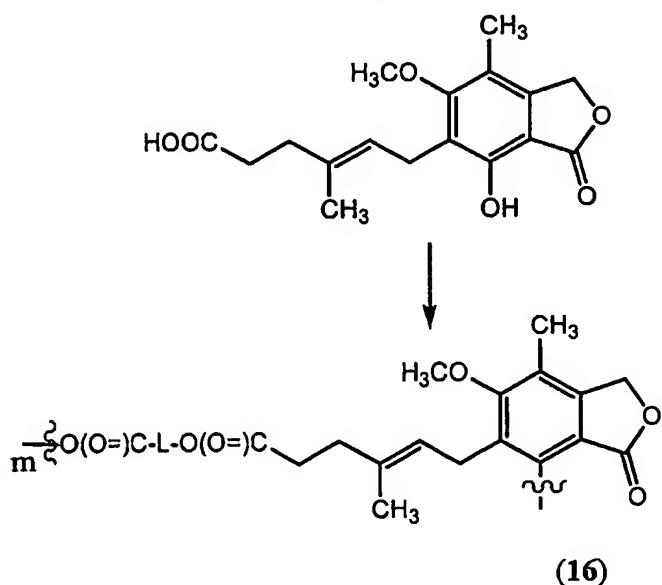
In the reaction illustrated in Scheme XIV, the linking group L is preferably  $-(CH_2)_x-$ , and more preferably, L is  $-(CH_2)_8-$ . A polymer of formula (15) wherein L has any of the values, specific values, or preferred values described herein is a preferred polymer of the invention. For a polymer of formula (15), m is an integer that is greater than or equal to 2. Prior to the polymerization illustrated in Scheme XIV, the carboxylic acid can be protected with a suitable protecting group, which can subsequently be removed, to provide the polymer of the invention.

5-Aminosalicilic acid can also be incorporated into a polymer of the invention which is a polymer of formula (23) or (24):



The preparation of a polymer of the invention comprising mycophenolic acid is illustrated in Scheme XV.

**Scheme XV**



5 In the reaction illustrated in Scheme XV, the linking group L is preferably  
- $(\text{CH}_2)_x$ -, and more preferably, L is - $(\text{CH}_2)_8$ - . A polymer of formula (16)  
wherein L has any of the values, specific values, or preferred values described  
herein is a preferred polymer of the invention. For a polymer of formula (16), m  
10 is an integer that is greater than or equal to 2.

### Activity

The ability of a polymer of the invention to produce a given therapeutic effect can be determined using *in vitro* and *in vivo* pharmacological models which are well known to the art.

All publications, patents, and patent documents (including the entire contents of U.S. Provisional Patent Application Number 60/220,707, filed 27 July 2000) are incorporated by reference herein, as though individually incorporated by reference. The invention has been described with reference to 5 various specific and preferred embodiments and techniques. However, it should be understood that many variations and modifications may be made while remaining within the spirit and scope of the invention.

**What is Claimed is:**

1. A polymer comprising a backbone, wherein the backbone comprises ester, thioester, or amide linkages, and wherein the backbone comprises one or more groups that will yield a biologically active compound upon hydrolysis of the polymer.
- 5
2. The polymer of claim 1 which comprises one or more units of formula (I) in the backbone:

10



wherein

$R_1$  is group that will yield a biologically active compound upon hydrolysis of the polymer;

15 each A is independently an ester linkage, a thioester linkage, or an amide linkage; and

L is a linking group.

- 20 3. The polymer of claim 1 or 2 wherein the biologically active compound is a non-steroidal ant-inflammatory drug, anti-bacterial, anti-fungal, anti-cancer, anti-thrombotic, immunosuppressive, or analgesic.
4. The polymer of claim 1 or 2 wherein the biologically active compound is atorvastatin; enalapril; ranitidine; ciprofloxacin; pravastatin; clarithromycin; cyclosporin; famotidine; leuprolide; acyclovir; paclitaxel; azithromycin; lamivudine; budesonide; albuterol; indinavir; metformin; alendronate; nizatidine; zidovudine; carboplatin; metoprolol; amoxicillin; diclofenac; lisinopril; ceftriaxone; captopril; salmeterol; imipenem; cilastatin; benazepril; cefaclor; ceftazidime; fluvastatin; quinapril; streptozocin; doxorubicin; daunorubicin; plicamycin; idarubicin; mitomycin C; pentostatin; mitoxantrone; cytarabine; fludarabine phosphate; floxuridine; cladribine; 6-mercaptopurine; thioguanine;
- 25
- 30

capecitabine; paclitaxel; docetaxel; etoposide; gemcitabine; topotecan; 5  
vinorelbine; vincristine; vinblastine; teniposide; melphalan; 2-p-  
sulfanilylanilinoethanol; 4,4'-sulfinyldianiline; 4-sulfanilamidosalicylic  
acid; acediasulfone; acetosulfone; amikacin; amphotericin B; ampicillin;  
10  
apalcillin; apicycline; apramycin; arbekacin; aspoxicillin; azidamfenicol;  
aztreonam; bacitracin; bambermycin(s); biapenem; brodimoprim;  
butirosin; capreomycin; carbenicillin; carbomycin; carumonam;  
cefadroxil; cefamandole; cefatrizine; cefbuperazone; cefclidin; cefdinir;  
15  
cefditoren; cefepime; cefetamet; cefixime; cefmenoxime; cefminox;  
cefodizime; cefonicid; cefoperazone; ceforanide; cefotaxime; cefotetan;  
cefotiam; cefozopran; cefpimizole; cefpiramide; cefpirome; cefprozil;  
cefroxadine; cesteram; ceftibuten; cefuzonam; cephalexin; cephaloglycin;  
20  
cephalosporin C; cephradine; chloramphenicol; chlortetracycline;  
clinafloxacin; clindamycin; clomocycline; colistin; cyclacillin; dapsone;  
demeclocycline; diathymosulfone; dibekacin; dihydrostreptomycin;  
dirithromycin; doxycycline; enoxacin; eniomyein; epicillin;  
25  
erythromycin; flomoxef; fortimicin(s); gentamicin(s); glucosulfone  
solasulfone; gramicidin S; gramicidin(s); grepafloxacin; guamecycline;  
hetacillin; isepamicin; josamycin; kanamycin(s); leucomycin(s);  
lincomycin; lomefloxacin; lucensomycin; lymecycline; mecloclcline;  
30  
meropenem; methacycline; micromomicin; midecamycin(s); minocycline;  
moxalactam; mupirocin; nadifloxacin; natamycin; neomycin; netilmicin;  
norfloxacin; oleandomycin; oxytetracycline; p-sulfanilylbenzylamine;  
panipenem; paromomycin; pazufloxacin; penicillin N; pipacycline;  
pipemidic acid; polymyxin; primycin; quinacillin; ribostamycin;  
rifamide; rifampin; rifamycin SV; rifapentine; rifaximin; ristocetin;  
ritipenem; rokitamycin; rolitetracycline; rosaramycin; roxithromycin;  
salazosulfadimidine; sancycline; sisomicin; sparfloxacin; spectinomycin;  
spiramycin; streptomycin; succisulfone; sulfachrysoidine; sulfaloxic  
acid; sulfamidochrysoidine; sulfanilic acid; sulfoxone; teicoplanin;  
temafloxacin; temocillin; tetroxoprim; thiamphenicol; thiazolsulfone;  
thiostrepton; ticarcillin; tigemonam; tobramycin; tosufloxacin;  
trimethoprim; trospectomycin; trovafloxacin; tuberactinomycin;

vancomycin; azaserine; candicidin(s); chlorphenesin; dermostatin(s); filipin; fungichromin; mepartycin; nystatin; oligomycin(s); perimycin A; tubercidin; 6-azaauridine; 6-diazo-5-oxo-L-norleucine; aclacinomycin(s); ancitabine; anthramycin; azacitidine; azaserine; bleomycin(s); carubicin; 5 carzinophillin A; chlorozotocin; chromomycin(s); denopterin; doxifluridine; edatrexate; eflornithine; elliptinium; enocitabine; epirubicin; mannomustine; menogaril; mitobronitol; mitolactol; mopidamol; nogalamycin; olivomycin(s); peplomycin; pirarubicin; piritrexim; prednimustine; procarbazine; pteropterin; puromycin; ranimustine; streptonigrin; thiamiprime; N-[[5-[(1,4-Dihydro-2-methyl-10 4-oxo-6-quinazolinyl)methyl]methylamino]-2-thienyl]carbonyl]-L-glutamic acid; trimetrexate, tubercidin, ubenimex, vindesine, zorubicin; argatroban; coumetarol; dicoumarol; ethyl biscoumacetate; ethylidene dicoumarol; iloprost; lamifiban; taprostene; tioclomarol; tirofiban; 15 amiprilose; bucillamine; gusperimus; oxymorphone; buprenorphine; butorphanol; nalbuphine; mycophenolic acid; procodazole; romurtide; sirolimus (rapamycin); tacrolimus; butethamine; fenalcomine; hydroxytetracaine; naepaine; orthocaine; piridocaine; salicyl alcohol; 3-amino-4-hydroxybutyric acid; aceclofenac; alminoprofen; amfenac; 20 bromfenac; bromosaligenin; bumadizon; carprofen; diclofenac; diflunisal; ditazol; enfenamic acid; etodolac; etofenamate; morphine; dopamine; bialamicol; fendosal; fepradinol; flufenamic acid; gentisic acid; glucamethacin; glycol salicylate; meclofenamic acid; mefenamic acid; mesalamine; niflumic acid; olsalazine; oxaceprol; S-adenosylmethionine; 25 salicylic acid; salsalate; sulfasalazine; tolfenamic acid; or methotrexate.

5. The polymer of claim 4 wherein the biologically active compound is morphine, dopamine, bialamicol, or tetracycline.
- 30 6. The polymer of claim 3 wherein the analgesic is oxymorphone, buprenorphine, butorphanol, nalbuphine, butethamine, fenalcomine, hydroxytetracaine, naepaine, orthocaine, piridocaine or salicyl alcohol.

7. The polymer of claim 3 wherein the anti-cancer compound is 6-azaauridine, 6-diazo-5-oxo-L-norleucine, 6-mercaptopurine, aclacinomycin(s), ancitabine, anthramycin, azacitidine, azaserine, bleomycin(s), capecitabine, carubicin, carzinophillin A, chlorozotocin, chromomycin(s), cladribine, cytarabine, daunorubicin, denopterin, docetaxel, doxifluridine, doxorubicin, edatrexate, eflornithine, elliptinium, enocitabine, epirubicin, etoposide, floxuridine, fludarabine, gemcitabine, idarubicin, mannomustine, melphalan, menogaril, methotrexate, mitobronitol, mitolactol, mitomycin C, mitoxantrone, 10 mopidamol, mycophenolic acid, nogalamycin, olivomycin(s), paclitaxel, pentostatin, peplomycin, pirarubicin, piritrexim, plicamycin, podophyllinic acid 2-ethylhydrazine, prednimustine, procarbazine, pteropterin, puromycin, ranimustine, streptonigrin, streptozocin, teniposide, thiamiprime, thioguanine, N-[[5-[[[(1,4-Dihydro-2-methyl-4-oxo-6-quinazolinyl)methyl]methylamino]-2-thienyl]carbonyl]-L- 15 glutamic acid, toptecan, trimetrexate, tubercidin, ubenimex, vinblastine, vindesine, vinorelbine or zorubicin .

8. The polymer of claim 3 wherein the nonsteroidal anti-inflammatory drug is 3-amino-4-hydroxybutyric acid, aceclofenac, alminoprofen, amfenac, 20 bromfenac, bromosaligenin, bumadizon, carprofen, diclofenac, diflunisal, ditazol, enfenamic acid, etodolac, etofenamate, fendosal, fepradinol, flufenamic acid, gentisic acid, glucamethacin, glycol salicylate, meclofenamic acid, mefenamic acid, mesalamine, niflumic acid, 25 olsalazine, oxaceprol, S-adenosylmethionine, salicylic acid, salsalate, sulfasalazine or tolfenamic acid.

9. The polymer of claim 3 wherein the anti-bacterial is 2-p- 30 sulfanilylanilinoethanol, 4,4'-sulfinyldianiline, 4-sulfanilamidosalicylic acid, acediasulfone, acetosulfone, amikacin, amoxicillin, amphotericin B, ampicillin, apalciliin, apicycline, apramycin, arbekacin, aspoxicillin, azidamfenicol, azithromycin, aztreonam, bacitracin, bambermycin(s), biapenem, brodimoprim, butirosin, capreomycin, carbenicillin,

carbomycin, carumonam, cefadroxil, cefamandole, cefatrizine,  
cefbuperazone, cefclidin, cefdinir, cefditoren, cefepime, cefetamet,  
cefixime, cefmenoxime, cefminox, cefodizime, cefonicid, cefoperazone,  
5 ceforanide, cefotaxime, cefotetan, cefotiam, cefozopran, cefpimizole,  
cefpipamide, cefpirome, cefprozil, cefroxadine, ceftazidime, cefteram,  
ceftibuten, ceftriaxone, cefuzonam, cephalexin, cephaloglycin,  
cephalosporin C, cephadrine, chloramphenicol, chlortetracycline,  
10 ciprofloxacin, clarithromycin, clinafloxacin, clindamycin, clomocycline,  
colistin, cyclacillin, dapsone, demeclocycline, diathymosulfone,  
dibekacin, dihydrostreptomycin, dirithromycin, doxycycline, enoxacin,  
enviomycin, epicillin, erythromycin, flomoxef, fortimicin(s),  
15 gentamicin(s), glucosulfone solasulfone, gramicidin S, gramicidin(s),  
grepafloxacin, guamecycline, hetacillin, imipenem, isepamicin,  
josamycin, kanamycin(s), leucomycin(s), lincomycin, lomefloxacin,  
lucensomycin, lymecycline, mectocycline, meropenem, methacycline,  
15 micronomicin, midecamycin(s), minocycline, moxalactam, mupirocin,  
nadifloxacin, natamycin, neomycin, netilmicin, norfloxacin,  
oleandomycin, oxytetracycline, p-sulfanilylbenzylamine, panipenem,  
paromomycin, pazufloxacin, penicillin N, pipacycline, pipemidic acid,  
20 polymyxin, primycin, quinacillin, ribostamycin, rifamide, rifampin,  
rifamycin SV, rifapentine, rifaximin, ristocetin, ritipenem, rokitamycin,  
rolitetracycline, rosaramycin, roxithromycin, salazosulfadimidine,  
sancycline, sisomicin, sparfloxacin, spectinomycin, spiramycin,  
streptomycin, succisulfone, sulfachrysoidine, sulfaloxic acid,  
25 sulfamidochrysoidine, sulfanilic acid, sulfoxone, teicoplanin,  
temafloxacin, temocillin, tetracycline, tetroxoprim, thiamphenicol,  
thiazolsulfone, thiostrepton, ticarcillin, tigemonam, tobramycin,  
tosufloxacin, trimethoprim, trospectomycin, trovafloxacin,  
tuberactinomycin or vancomycin.

30

10. The polymer of claim 3 wherein the anti-fungal is azaserine,  
candidin(s), chlorphenesin, dermostatin(s), filipin, fungichromin,  
meparticin, nystatin, oligomycin(s), perimycin A or tubercidin.

11. The polymer of claim 3 wherein the anti-thrombotic is argatroban, coumetarol, dicoumarol, ethyl biscoumacetate, ethylidene dicoumarol, iloprost, lamifiban, taprostene, tioclomarol or tirofiban.
- 5
12. The polymer of claim 3 wherein the immunosuppressive is 6-mercaptopurine, amiprilose, bucillamine, gusperimus, mycophenolic acid, procodazole, romurtide, sirolimus (rapamycin), tacrolimus or ubenimex
- 10
13. The polymer of claim 4 wherein the biologically active compound is amikacin; amphotericin B; apicycline; apramycin; arbekacin; azidamfenicol; bambermycin(s); butirosin, carbomycin; cefpiramide; chloramphenicol; chlortetracycline; clindamycin; clomocycline; demeclocycline; diathymosulfone; dibekacin, dihydrostreptomycin; dirithromycin; doxycycline; erythromycin; fortimicin(s); gentamicin(s); glucosulfone solasulfone; guamecycline; isepamicin; josamycin; kanamycin(s); leucomycin(s); lincomycin; lucensomycin; lymecycline; mecloxycline; methacycline; micronomicin; midecamycin(s); minocycline; mupirocin; natamycin; neomycin; netilmicin; oleandomycin; oxytetracycline; paromomycin; pipacycline; podophyllinic acid 2-ethylhydrazine; primycin; ribostamycin; rifamide; rifampin; rifamycin SV; rifapentine; rifaximin; ristocetin; rokitamycin; rolitetracycline; rosaramycin; roxithromycin; sancycline; sisomicin; spectinomycin; spiramycin; streptomycin; teicoplanin; tetracycline; thiamphenicol; thiostrepton; tobramycin; trospectomycin; tuberactinomycin; vancomycin; candididin(s); chlorphenesin; dermostatin(s); filipin; fungichromin; meparticin; nystatin; oligomycin(s); perimycin A; tubercidin; 6-azauridine; aclacinomycin(s); ancitabine; anthramycin; azacitidine; bleomycin(s); carubicin; carzinophillin A; chlorozotocin; chromomycin(s); doxifluridine; enocitabine; epirubicin; gemcitabine; mannomustine; menogaril; atorvastatin; pravastatin; clarithromycin; leuprolide; paclitaxel;
- 15
- 20
- 25
- 30

mitobronitol; mitolactol; molidamol; nogalamycin; olivomycin(s);  
peplomycin; pirarubicin; prednimustine; puromycin; ranimustine;  
tubercidin; vindesine; zorubicin; coumetarol; dicoumarol; ethyl  
5 biscoumacetate; ethylidene dicoumarol; iloprost; taprostene; tioclofarol;  
amiprilose; romurtide; sirolimus (rapamycin); tacrolimus; salicyl alcohol;  
bromosaligenin; ditazol; fepradolin; gentisic acid; glucamethacin;  
olsalazine; S-adenosylmethionine; azithromycin; salmeterol; budesonide;  
albuterol; indinavir; fluvastatin; streptozocin; doxorubicin; daunorubicin;  
10 plicamycin; idarubicin; pentostatin; mitoxantrone; cytarabine;  
fludarabine phosphate; floxuridine; cladribine; capecitabine; docetaxel;  
etoposide; topotecan; vinblastine; or teniposide.

14. The polymer of claim 4 wherein the biologically active compound is 2-p-sulfanilylanilinoethanol; 4,4'-sulfinyldianiline; acediasulfone;

15 acetosulfone; amikacin; apramycin; arbekacin; bacitracin; brodimoprim;  
butirosin; colistin; capreomycin; dapsone; dibekacin; enviomycin;  
glucosulfone solasulfone; gramicidin S; polymyxin; teicoplanin;  
fortimicin(s); gentamicin(s); grepafloxacin; imipenem; isepamicin;  
kanamycin(s); lymecycline; micromomicin; neomycin; netilmicin; p-

20 sulfanilylbenzylamine; paromomycin; ribostamycin; ristocetin;  
sisomicin; sparfloxacin; spectinomycin; sulfachrysoidine;

sulfamidochrysoidine; sulfoxone; tetroxoprim; thiazolsulfone;  
tobramycin; trimethoprim; edatrexate; eflornithine; mannomustine;  
mitoxantrone; peplomycin; piritrexim; procarbazine; pteropterin;

25 trimetrexate; gusperimus; butethamine; naepaine; piridocaine;  
trospectomycin; tuberactinomycin; vancomycin; candidin(s);  
mepartycin; perimycin A; ranitidine; famotidine; metformin; nizatidine;  
carboplatin; lisinopril; methotrexate; mitomycin; bleomycin(s); or  
thioguanine.

30

15. The polymer of claim 4 wherein the biologically active compound is 2-p-sulfanilylanilinoethanol; 4-sulfanilamidosalicylic acid; amikacin;  
amphotericin B; apramycin; arbekacin; aspoxicillin; butirosin;

capreomycin; cefadroxil; cefatrizine; cefdinir; cefprozil; dibekacin; dihydrostreptomycin; dirithromycin; enviomycin; gramicidin(s); teicoplanin; fortimicin(s); gentamicin(s); glucosulfone solasulfone; isepamicin; kanamycin(s); licensomycin; lymecycline; meropenem; 5 micronomicin; natamycin; neomycin; netilmicin; paromomycin; ribostamycin; ristocetin; sisomicin; spectinomycin; streptomycin; thiostrepton; tobramycin; trospectomycin; tuberactinomycin; vancomycin; candicidin(s); mepartycin; nystatin; perimycin A; tubercidin; anthramycin; azacitidine; bleomycin(s); carubicin; 10 carzinophillin A; cytarabine; denopterin; elliptinium; epirubicin; mannomustine; peplomycin; pirarubicin; pteropterin; puromycin; streptonigrin; tubercidin; ubenimex; vindesine; zorubicin; gusperimus; ubenimex; fenalcomine; hydroxytetracaine; orthocaine; 3-amino-4- 15 hydroxybutyric acid; etofenamate; fepradolin; mesalamine; S-adenosylmethionine; leuprolide; acyclovir; paclitaxel; lamivudine; albuterol; indinavir; alendronate; zidovudine; metoprolol; amoxicillin; salmeterol; imipenem; doxorubicin; daunorubicin; idarubicin; 20 pentostatin; mitoxantrone; fludarabine phosphate; floxuridine; cladribine; vinorelbine; vincristine; or vinblastine.

25 16. The polymer of claim 4 wherein the biologically active compound is bambermycin(s); carbenicillin; carzinophillin A; cefixime; cefminox; cefpimizole; cefodizime; cefonicid; ceforanide; cefotetan; cefributene; cephalosporin C; denopterin; edatrexate; moxalactam; olsalazine; penicillin N; quinacillin; temocillin; ticarcillin; N-[[5-[(1,4-Dihydro-2- 30 methyl-4-oxo-6-quinazolinyl)methyl]methylamino]-2-thienyl]carbonyl]-L-glutamic acid; lisinopril; cilastatin; ceftazidime; or methotrexate.

35 17. The polymer of claim 4 wherein the biologically active compound is 4-sulfanilamidosalicylic acid; amphotericin B; apalcillin; apicycline; aspoxicillin; bambermycin(s); biapenem; cefadroxil; cefamandole; cefatrizine; cefbuperazone; cefdinir; cefonicid; cefoperazone; cefpiramide; cefprozil; enviomycin; teicoplanin; flomoxef; glycol

salicylate; lucensomycin; lymecycline; meropenem; moxalactam; mupirocin; nadifloxacin; natamycin; panipenem; podophyllinic acid 2-ethylhydrazine; ritipenem; salazosulfadimidine; sulfaloxic acid; vancomycin; 3-amino-4-hydroxybutyric acid; candididin(s); carzinophillin A; denopterin; diflunisal; fendosal; gentisic acid; iloprost; lamifiban; mesalamine; nystatin; olsalazine; oxaceprol; pteropterin; romurtide; salicylic acid; salsalate; streptonigrin; sulfasalazine; taprostene; ubenimex; amoxicillin; pravastatin; xinafoate; imipenem; mycophenolic acid; or fluvastatin.

10

18. The polymer of claim 4 wherein the biologically active compound is 3-amino-4-hydroxybutyric acid; 4-sulfanilamidosalicylic acid; 6-diazo-5-oxo-L-norleucine; aceclofenac; acediasulfone; alminoprofen; amfenac; amphotericin B; ampicillin; argatroban; aspoxicillin; azaserine; aztreonam; bromfenac; bumadizon; candididin(s); carprofen; carumonam; carzinophillin A; cefadroxil; cefatrizine; cefclidin; cefdinir; cefditoren; cefepime; cefetamet; cefixime; cefmenoxime; cefminox; cefodizime; ceforanide; cefotaxime; cefotiam; cefozopran; cefpirome; cefprozil; cefroxadine; ceftazidime; cefteram; cefributen; ceftriaxone; cefuzonam; cephalexin; cephaloglycin; cephalosporin C; cephadrine; clinafloxacin; cyclacillin; denopterin; edatrexate; eflornithine; enfenamic acid; enoxacin; epicillin; etodolac; eniomycin; teicoplanin; flufenamic acid; grepafloxacin, hetacillin; imipenem; lomefloxacin; lucensomycin; lymecycline; meclofenamic acid; mefenamic acid; meropenem; mesalamine; natamycin; niflumic acid; norfloxacin; nystatin; pazufloxacin; penicillin N; pipemidic acid; procodazole; pteropterin; S-adenosylmethionine; sparfloxacin; streptonigrin; succisulfone; sulfachrysoidine; temafloxacin; tigemonam; tirofiban; tolfenamic acid; tosufloxacin; trovafloxacin; ubenimex; vancomycin; enalapril; amoxicillin; ciprofloxacin; diclofenac; lisinopril; ceftriaxone; cilastatin; benazepril; cefaclor; ceftazidime; quinapril; melphalan; or methotrexate.

19. The polymer of claim 4 wherein the biologically active compound is phenamidine, acriflavine, chloroazodin, arsphenamine, amicarbilide or aminoquinuride.

5 20. The polymer of claim 2 which is a polymer of formula (7), (8), (9), (10), (11), (12), (13), (14), (15), or (16), or a polymer of formula (III), (IV), (V), or (VI), as illustrated herein above.

21. The polymer of claim 1 which comprises one or more units of formula 10 (II) in the backbone:



wherein

15  $\text{R}_2$  and  $\text{R}_3$  are each independently a group that will yield a biologically active compound upon hydrolysis of the polymer; each A is independently an amide or ester linkage; and each L is independently a linking group.

20 22. The polymer of claim 2 wherein L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 1 to 25 carbon atoms, wherein one or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-), and wherein the chain is optionally substituted on carbon with one or more (e.g. 1, 2, 3, or 4) substituents selected from the group consisting of  $(\text{C}_1\text{-C}_6)$ alkoxy,  $(\text{C}_3\text{-C}_6)$ cycloalkyl,  $(\text{C}_1\text{-C}_6)$ alkanoyl,  $(\text{C}_1\text{-C}_6)$ alkanoyloxy,  $(\text{C}_1\text{-C}_6)$ alkoxycarbonyl,  $(\text{C}_1\text{-C}_6)$ alkylthio, azido, cyano, nitro, halo, hydroxy, oxo, carboxy, aryl, aryloxy, heteroaryl, and heteroaryloxy.

25 30 23. The polymer of claim 2 wherein L is a divalent, branched or unbranched, saturated or unsaturated, hydrocarbon chain, having from 1 to 25 carbon atoms, wherein the chain is optionally substituted on carbon with one or more (e.g. 1, 2, 3, or 4) substituents selected from the group consisting of

(C<sub>1</sub>-C<sub>6</sub>)alkoxy, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyloxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxycarbonyl, (C<sub>1</sub>-C<sub>6</sub>)alkylthio, azido, cyano, nitro, halo, hydroxy, oxo, carboxy, aryl, aryloxy, heteroaryl, and heteroaryloxy.

5 24. The polymer of claim 2 wherein L is a peptide.

25. The polymer of claim 2 wherein L is an amino acid.

26. The polymer of claim 2 wherein L is a divalent, branched or unbranched, 10 saturated or unsaturated, hydrocarbon chain, having from 1 to 25 carbon atoms, wherein one or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-).

27. The polymer of claim 2 wherein L is a divalent, branched or unbranched, 15 saturated or unsaturated, hydrocarbon chain, having from 3 to 15 carbon atoms, wherein one or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-), and wherein the chain is optionally substituted on carbon with one or more (e.g. 1, 2, 3, or 4) substituents selected from the group consisting of (C<sub>1</sub>-C<sub>6</sub>)alkoxy, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyl, (C<sub>1</sub>-C<sub>6</sub>)alkanoyloxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxycarbonyl, (C<sub>1</sub>-C<sub>6</sub>)alkylthio, azido, cyano, nitro, halo, hydroxy, oxo, carboxy, aryl, aryloxy, heteroaryl, and heteroaryloxy.

28. The polymer of claim 2 wherein L is a divalent, branched or unbranched, 25 saturated or unsaturated, hydrocarbon chain, having from 3 to 15 carbon atoms, wherein one or more (e.g. 1, 2, 3, or 4) of the carbon atoms is optionally replaced by (-O-) or (-NR-).

29. The polymer of claim 2 wherein L is a divalent, branched or unbranched, 30 saturated or unsaturated, hydrocarbon chain, having from 3 to 15 carbon atoms.

30. The polymer of claim 2 wherein L is a divalent, branched or unbranched, hydrocarbon chain, having from 3 to 15 carbon atoms.
31. The polymer of claim 2 wherein L is a divalent, branched or unbranched, hydrocarbon chain, having from 6 to 10 carbon atoms.
- 5 32. The polymer of claim 2 wherein L is a divalent hydrocarbon chain having 7, 8, or 9 carbon atoms.
- 10 33. The polymer of claim 2 wherein L is a divalent hydrocarbon chain having 8 carbon atoms.
34. A biocompatible and biodegradable polyester or polyamide comprising a biologically active compound containing at least two alcohol or phenol groups or at least two amine groups co-polymerized to bis(acyl) chlorides or carboxylic acids.
- 15 35. The polymer of claim 1 further comprising another therapeutic agent dispersed in the matrix of the polymer.
- 20 36. The polymer of claim 1 further comprising another therapeutic agent appended to the polymer backbone.
37. A pharmaceutical composition comprising a polymer of claim 1 and a pharmaceutically acceptable carrier.
- 25 38. A therapeutic method for treating a disease in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 1.
- 30 39. A therapeutic method for producing an analgesic effect in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 6.

40. A therapeutic method for treating cancer comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 7.
- 5 41. A therapeutic method for producing an anti-inflammatory effect in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 8.
- 10 42. A therapeutic method for producing an anti-bacterial effect in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 9.
- 15 43. A therapeutic method for producing an anti-fungal effect in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 10.
- 20 44. A therapeutic method for producing an immunosuppressive effect in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 12.
45. A therapeutic method for producing an anti-thrombotic effect in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of claim 11.
- 25 46. A method for producing a biocompatible and biodegradable polyester or polyamide which degrades into a biologically active compound comprising co-polymerizing a biologically active compound containing at least two alcohol or phenol groups or at least two amine groups with carboxylic acid groups or bis(acyl) chlorides.
- 30 47. A method of delivering a biologically active compound to a host comprising administering to the host a biocompatible and biodegradable polyester or polyamide of claim 1.

48. A therapeutic method for treating psoriasis, inflammatory bowel disease, skin cancer, or a brain tumor comprising administering to a mammal in need of such therapy, an effective amount of a polymer of any one of formulae 10, 11, 12 or 20 as described herein.

5

49. A therapeutic method for producing an anti-infective effect in an animal comprising administering to an animal in need of such therapy, an effective amount of a polymer of any one of formulae 10, 11, 12 or 20 as described herein.

10

50. A therapeutic method for treating cancer comprising administering to an animal in need of such therapy, an effective amount of a polymer of any one of formulae 10, 11, 12, 13, 14 or 20 as described herein.

15

51. A therapeutic method for treating pain comprising administering to an animal in need of such therapy, an effective amount of a polymer of formula 7 as described herein.

52. A therapeutic method for treating Parkinson's disease comprising

20

administering to a mammal in need of such therapy, an effective amount of a polymer of formula 8 as described herein.

53. The polymer as described in claim 1 for use in medical therapy.

25

54. The use of a polymer as described in claim 1 for the manufacture of a medicament useful for the treatment of a disease in a mammal, such as a human.